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COMPOUNDS AND METHODS FOR MODULATING CEREBRAL AMYLOID ANGIOPATHY

RELATED APPLICATIONS

This application claims the benefit of priority under 35 U.S.C. 119(e) to copending U.S. 5 Provisional Application No. 60/171,877, filed December 23, 1999, the entire contents of which are incorporated herein by reference.

BACKGROUND OF THE INVENTION

Cerebral amyloid angiopathy (CAA) remains a largely untreatable disease often not diagnosed until autopsy. It ranges in severity from asymptomatic amyloid deposition in 10 otherwise normal cerebral vessels to complete replacement and breakdown of the cerebrovascular wall. Severe CAA can cause lobar cerebral hemorrhage, transient neurologic symptoms, and dementia with leukoencephalopathy. (see Greenberg, *Neurology* 1998, 51:690-694).

Amyloid- β (A β) is a toxic peptide which is implicated in the pathogenesis of CAA. A β 15 peptide is derived from a normal proteolytic cleavage of the precursor protein, the Amyloid- β precursor protein (β APP). Advanced cases of CAA demonstrate structural changes to the walls of the amyloid-laden vessel such as cracking between layers, smooth muscle cell toxicity, microaneurysm formation, and fibrinoid necrosis.

The exact mechanisms involved in the genesis of cerebral amyloid angiopathy (CAA) 20 have not been completely established, but it appears that a preponderance of the form of the 39-40 amino acid A β peptide (A β 40) is responsible for the deposits on blood vessel wall cells which lead to CAA, in comparison to the 42-43 amino acid A β peptides (A β 42 and A β 43), which are implicated in other amyloid-related conditions such as Alzheimer's Disease (AD).

SUMMARY OF THE INVENTION

25 The present invention provides methods for modulating, e.g., inhibiting and/or preventing, cerebral amyloid angiopathy. The present invention is based, at least in part, on the discovery that compounds which interfere with the deposition of A β peptide, e.g., the A β 40 peptide, in blood vessel wall cells, prevent the structural changes to cerebral blood vessels like capillaries, that lead to CAA. It is believed, without intending to limit the invention as claimed 30 herein, that the compounds of the invention interfere with the association of the A β 40 peptide,

e.g., the association of the A β 40 peptide to the sulfate GAGs present at the smooth muscle cell surface, and thus prevent intracellular and extracellular amyloid deposition. However, while it is believed that inhibition of A β 40 is a significant factor in inhibiting CAA, the A β 40 inhibitors of the invention may well work in other ways to inhibit or prevent CAA, and these are intended 5 to be part of the present invention.

Accordingly, this invention pertains to a method of modulating, e.g., inhibiting and/or preventing, cerebral amyloid angiopathy. The method includes contacting a blood vessel wall cell with an A β 40 inhibitor, such that the compound inhibits or prevents cerebral amyloid angiopathy. The A β 40 inhibitor is believed to at least interfere with the ability of the A β 40 10 peptide to form amyloid fibrils and/or with the ability of the A β 40 peptide to bind to a cell (e.g., blood vessel wall smooth muscle cells, pericytes or endothelial cells) surface molecule or structure, forming deposits on the walls of the blood vessel and thus prevent A β -induced cell death and/or the structural changes to cerebral blood vessels, e.g., capillaries, medium sized arteries, or arterioles, that lead to CAA. The A β 40 peptide can be either in a soluble form or in 15 a fibril form.

In one embodiment, the A β 40 inhibitor may be ethanesulfonic acid, 1,2-ethanedisulfonic acid, 1-propanesulfonic acid, 1,3-propanedisulfonic acid, 1,4-butanedisulfonic acid, 1,5-pantanedisulfonic acid, 2-aminoethanesulfonic acid, or 4-hydroxy-1-butanesulfonic acid, and pharmaceutically acceptable salts thereof. In other preferred embodiments, the A β 40 20 inhibitor may be 1-butanesulfonic acid, 1-decanesulfonic acid, 2-propanesulfonic acid, 3-pentanesulfonic acid, or 4-heptanesulfonic acid, and pharmaceutically acceptable salts thereof. In yet further preferred embodiments, the A β 40 inhibitor may be 1,7-dihydroxy-4-heptanesulfonic acid, 3-amino-1-propanesulfonic acid, or a pharmaceutically acceptable salt thereof. In another embodiment the A β 40 inhibitor is a peptide or a peptidomimetic which 25 interacts with specific regions of the A β peptide such as the regions responsible for cellular adherence (aa 10-16), GAG binding site region (13-16) or the region responsible for the β -sheet formation (16-21). These peptides are the d-stereoisomers of the A β or complementary image of the A β peptide.

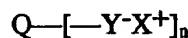
In one embodiment, the A β 40 inhibitor is administered in a pharmaceutically acceptable 30 formulation. The pharmaceutically acceptable formulation can be a dispersion system like a

lipid-based formulation, a liposome formulation, or a multivesicular liposome formulation. The pharmaceutically acceptable formulation can also comprise a polymeric matrix, e.g., synthetic polymers such as polyesters (PLA, PLGA), polyethylene glycol, poloxomers, polyanhydrides, and pluronic; or naturally derived polymers, such as albumin, alginate, cellulose derivatives, collagen, fibrin, gelatin, and polysaccharides. In other preferred embodiments, the pharmaceutically acceptable formulation provides sustained delivery of the A_β40 inhibitor to the target site.

Yet another aspect of the invention pertains to a method of treating a disease state characterized by cerebral amyloid angiopathy in a subject. The method includes administering an A_β40 inhibitor to the subject, such that the disease state characterized by cerebral amyloid angiopathy is treated, e.g., inhibited or prevented.

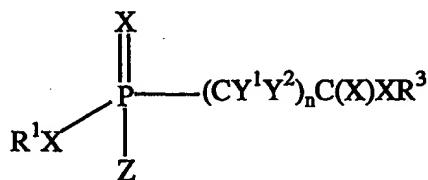
Another aspect of the invention pertains to a method of modulating, e.g., inhibiting and/or preventing, cerebral amyloid angiopathy, including contacting a blood vessel wall cell with an A_β40 inhibitor having the structure:

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wherein Y⁻ is an anionic group at physiological pH; Q is a carrier group; X⁺ is a cationic group; and n is an integer selected such that the biodistribution of the A_β40 inhibitor for an intended target site is not prevented while maintaining activity of the A_β40 inhibitor, provided that the A_β40 inhibitor is not chondroitin sulfate A, such that cerebral amyloid angiopathy is inhibited or prevented.

In yet another aspect, the invention features a method of modulating, e.g., inhibiting and/or preventing, cerebral amyloid angiopathy, including contacting a blood vessel wall cell with an A_β40 inhibitor having the structure:



25 wherein Z is XR² or R⁴, R¹ and R² are each independently hydrogen, a substituted or unsubstituted aliphatic group (preferably a branched or straight-chain aliphatic moiety having from 1 to 24 carbon atoms in the chain; or an unsubstituted or substituted cyclic aliphatic

moiety having from 4 to 7 carbon atoms in the aliphatic ring; preferred aliphatic and cyclic aliphatic groups are alkyl groups, more preferably lower alkyl), an aryl group, a heterocyclic group, or a salt-forming cation; R³ is hydrogen, lower alkyl, aryl, or a salt-forming cation; R⁴ is hydrogen, lower alkyl, aryl or amino (including alkylamino, dialkylamino (including cyclic 5 amino moieties), arylamino, diarylamino, and alkylarylamino); X is, independently for each occurrence, O or S; Y¹ and Y² are each independently hydrogen, halogen (e.g., F, Cl, Br, or I), alkyl (preferably lower alkyl), amino, hydroxy, alkoxy, or aryloxy; and n is an integer from 0 to 12 (more preferably 0 to 6, more preferably 0 or 1), such that cerebral amyloid angiopathy is inhibited or prevented.

10 Other features and advantages of the invention will be apparent from the following detailed description, and from the claims.

DETAILED DESCRIPTION OF THE INVENTION

The present invention is based, at least in part, on the discovery that compounds which interfere with the ability of the A β 40 peptide to form deposits in cerebral blood vessels, e.g., on 15 the smooth muscle cells thereof, and thus prevent the structural changes to cerebral blood vessels that lead to CAA.

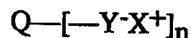
As used herein, the language "contacting" is intended to include both *in vivo*, *in vitro*, or *ex vivo* methods of bringing an A β 40 inhibitor into proximity with a blood vessel wall cell,

The term "blood vessel wall cell" includes smooth muscle cells, pericytes and endothelial cells. In a preferred embodiment the blood vessel wall cell is a smooth muscle cell.

A β 40 inhibitors

In one embodiment, the method of the invention includes contacting a blood vessel wall cell *in vitro* or administering to a subject *in vivo*, an effective amount of an A β 40 inhibitor, which has at least one anionic group covalently attached to a carrier molecule. As used herein, an "A β 40 inhibitor" includes compounds which can interfere with the ability of a CAA-associated A β peptide, e.g., A β 40, to either form fibrils or interact with a cell surface molecule such as a proteoglycan constituent of a basement membrane, e.g., a glycosaminoglycan. An A β 40 inhibitor can interfere with the ability of both fibrillar or non-fibrillar CAA-associated A β peptide, e.g., A β 40, to interact with a cell surface molecule.

The A β 40 inhibitor can have the structure:

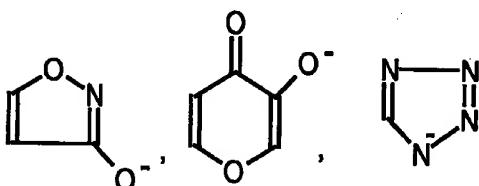


wherein Y⁻ is an anionic group at physiological pH; Q is a carrier group; X⁺ is a cationic group; and n is an integer. The number of anionic groups ("n") is selected such that the biodistribution of the A β 40 inhibitor for an intended target site is not prevented while maintaining activity of the A β 40 inhibitor. For example, the number of anionic groups is not so great as to prevent traversal of an anatomical barrier, such as a cell membrane, or entry across a physiological barrier, such as the blood-brain barrier. In one embodiment, n is an integer between 1 and 10. In another embodiment, n is an integer between 3 and 8. These compounds are described in U.S. Patent Nos. 5,643,562, 5,972,328, 5,728,375, 5,840,294, and U.S. Application No. 60/131,464. Such compounds also include or can be described as glycosaminoglycan ("GAG") mimics or mimetics. Other compounds which may be included are those described in, e.g., Pillot et al., *Eur. J. Biochem* vol. 243 No. 3, 1997 (apoE2, apoE3); WO98/22441; WO98/22430; WO96/10220; WO96/07425; and WO96/39834.

An anionic group of an A β 40 inhibitor of the invention is a negatively charged moiety that, when attached to a carrier group, can interfere with the ability of a CAA-associated A β peptide, e.g., A β 40, to either form fibrils or interact with a cell surface molecule such as a proteoglycan constituent of a basement membrane, e.g., a glycosaminoglycan ("GAG"). As such, A β 40 is inhibited from forming deposits in blood vessels, e.g., cerebral blood vessel wall

smooth muscle cells, thus preventing hardening of the vessel walls and, therefore, cerebral amyloid angiopathy.

For purposes of this invention, the anionic group is negatively charged at physiological pH. Preferably, the anionic A β 40 inhibitor mimics the structure of a sulfated proteoglycan, i.e., 5 is a sulfated compound or a functional equivalent thereof. "Functional equivalents" of sulfates are intended to include compounds such as sulfamates as well as bioisosteres. Bioisosteres encompass both classical bioisosteric equivalents and non-classical bioisosteric equivalents. Classical and non-classical bioisosteres of sulfate groups are known in the art (see e.g., Silverman, R.B. *The Organic Chemistry of Drug Design and Drug Action*, Academic Press, 10 Inc.: San Diego, CA, 1992, pp.19-23). Accordingly, an A β 40 inhibitor of the invention can comprise at least one anionic group including sulfonates, sulfates, sulfamates, phosphonates, phosphates, carboxylates, and heterocyclic groups of the following formulae:



Depending on the carrier group, more than one anionic group can be attached thereto.

15 When more than one anionic group is attached to a carrier group, the multiple anionic groups can be the same structural group (e.g., all sulfonates) or, alternatively, a combination of different anionic groups can be used (e.g., sulfonates, phosphonates, and sulfates, etc.).

The ability of an A β 40 inhibitor of the invention to inhibit an interaction between A β 40 peptide and a glycoprotein or proteoglycan constituent of a basement membrane can be 20 assessed by an *in vitro* binding assay, such as the one described in Leveugle B. et al. (1998) *J. of Neurochem.* 70(2):736-744. Briefly, a constituent of the basement membrane, preferably a glycosaminoglycan (GAG) can be radiolabeled, e.g., at a specific activity of 10,000 cpm, and then incubated with A β 40 peptide-Sepharose beads at, for example, a ratio of 5:1 (v/v) in the presence or absence of the A β 40 inhibitor. The A β 40 peptide-Sepharose beads and the 25 radiolabeled GAG can be incubated for approximately 30 minutes at room temperature and then the beads can be successively washed with a Tris buffer solution containing NaCl (0.55 M and 2 M). The binding of the basement membrane constituent (e.g., GAG) to the A β 40 peptide can then be measured by collecting the fractions from the washings and subjecting them to

scintillation counting. An A β 40 inhibitor which inhibits an interaction between A β 40 and a glycoprotein or proteoglycan constituent of a basement membrane, e.g., GAG, will increase the amount of radioactivity detected in the washings.

In the same manner, the invention relates to a method of diagnosing CAA *in vivo*,
5 whereas an labeled inhibitor of the invention is administered to a subject and the disposition of the inhibitor is determined to see whether a CAA-related condition exists. The label may be one conventionally known in the art which allows for detection of the compound either *in vivo* or *in vitro*, e.g., radiolabel, fluorescent, etc. Using techniques with which those of ordinary skill in the art will be familiar, e.g., PET scan, bound tagged inhibitor of the invention may be
10 visualized, e.g., in regions where CAA would be found, such as near the cerebellum.

Preferably, an A β 40 inhibitor of the invention interacts with a binding site for a basement membrane glycoprotein or proteoglycan in A β 40 and thereby inhibits the binding of the A β 40 peptide to the basement membrane constituent, e.g., GAG. Basement membrane glycoproteins and proteoglycans include GAG, laminin, collagen type IV, fibronectin,
15 chondroitin sulfate, perlecan, and heparan sulfate proteoglycan (HSPG). In a preferred embodiment, the therapeutic compound inhibits an interaction between an A β 40 peptide and a GAG. Consensus binding site motifs for GAG in amyloidogenic proteins have been described (see, for example, Hileman R. E. et al. (1998) *BioEssays* 20:156-167). For example, a GAG consensus binding motif can be of the general formula X-B-B-X-B-X or X-B-B-B-X-X-B-X,
20 wherein B are basic amino acids (e.g., lysine or arginine) and X are hydrophobic amino acids. A GAG consensus binding motif can further be of the general formula T-X-X-B-X-X-T-B-X-X-X-T-B-B, wherein T defines a turn of a basic amino acid, Bs are basic amino acids (e.g., lysine, arginine, or occasionally glutamine) and X are hydrophobic amino acids. The distance between the first and the second turn can range from approximately 12 Å to 17 Å. The distance between
25 the second and the third turn can be approximately 14 Å. The distance between the first and the third turn can range from approximately 13 Å to 18 Å.

Accordingly, in the A β 40 inhibitors of the invention, when multiple anionic groups are attached to a carrier group, the relative spacing of the anionic groups can be chosen such that the anionic groups (e.g., sulfonates or phosphonates) optimally interact with the basic residues
30 within the GAG binding site (thereby inhibiting interaction of GAG with the site). For example, anionic groups can be spaced approximately 15 ± 1.5 Å, 14 ± 1.5 Å and/or 16 ± 1.5 Å

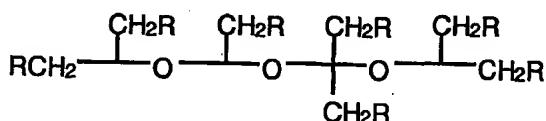
apart, or appropriate multiples thereof, such that the relative spacing of the anionic groups allows for optimal interaction with a binding site for a basement membrane constituent (e.g., GAG) in an A β 40 peptide.

A β 40 inhibitors of the invention typically further comprise a counter cation (i.e., X $^+$ in the general formula: Q—[—Y-X $^+$] $_n$). Cationic groups include positively charged atoms and moieties. If the cationic group is hydrogen, H $^+$, then the compound is considered an acid, e.g., ethanesulfonic acid. If hydrogen is replaced by a metal or its equivalent, the compound is a salt of the acid. Pharmaceutically acceptable salts of the A β 40 inhibitor are within the scope of the invention. For example, X $^+$ can be a pharmaceutically acceptable alkali metal, alkaline earth, higher valency cation, polycationic counter ion or ammonium. A preferred pharmaceutically acceptable salt is a sodium salt but other salts are also contemplated within their pharmaceutically acceptable range.

Within the A β 40 inhibitor, the anionic group(s) is covalently attached to a carrier group. Suitable carrier groups include aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups, and groups derived from carbohydrates, polymers, peptides, peptide derivatives, or combinations thereof. A carrier group can be substituted, e.g., with one or more amino, nitro, halogen, thiol or hydroxyl groups.

As used herein, the term "carbohydrate" is intended to include substituted and unsubstituted mono-, oligo- and polysaccharides. Monosaccharides are simple sugars usually

derived polymers and copolymers and salts thereof. In one embodiment, the polymer has a molecular weight of approximately 800-1000 Daltons. Examples of polymers with suitable covalently attached anionic groups (e.g., sulfonates or sulfates) include poly(2-acrylamido-2-methyl-1-propanesulfonic acid); poly(2-acrylamido-2-methyl-1-propanesulfonic acid-co-acrylonitrile); poly(2-acrylamido-2-methyl-1-propanesulfonic acid-co-styrene); poly(vinylsulfonic acid); poly(sodium 4-styrenesulfonic acid); and sulfates and/or sulfonates derived from: poly(acrylic acid); poly(methyl acrylate); poly(methyl methacrylate); and poly(vinyl alcohol); and pharmaceutically acceptable salts thereof. Examples of polymers with suitable covalently attached anionic groups include those of the formula:



wherein R is SO_3H or OSO_3H ; and pharmaceutically acceptable salts thereof.

Peptides and peptide derivatives can also act as carriers. The term "peptide" includes two or more amino acids covalently attached through a peptide bond. Amino acids which can be used in peptide carriers include those naturally occurring amino acids found in proteins such as glycine, alanine, valine, cysteine, leucine, isoleucine, serine, threonine, methionine, glutamic acid, aspartic acid, glutamine, asparagine, lysine, arginine, proline, histidine, phenylalanine, tyrosine, and tryptophan. The term "amino acid" further includes analogs, derivatives and congeners of naturally occurring amino acids, one or more of which can be present in a peptide derivative. For example, amino acid analogs can have lengthened or shortened side chains or variant side chains with appropriate functional groups. Also included are the D and L stereoisomers of an amino acid when the structure of the amino acid admits of stereoisomeric forms. The term "peptide derivative" further includes compounds which contain molecules which mimic a peptide backbone but are not amino acids (so-called peptidomimetics), such as benzodiazepine molecules (see e.g., James, G. L. et al. (1993) *Science* 260:1937-1942). The anionic groups can be attached to a peptide or peptide derivative through a functional group on the side chain of certain amino acids or other suitable functional group. For example, a sulfate group can be attached through the hydroxyl side chain of a serine residue. A peptide can be designed to interact with a binding site for a basement membrane constituent (e.g., a GAG) in an $\text{A}\beta 40$ peptide (as described above). Accordingly, in one embodiment, the peptide comprises

four amino acids and anionic groups (e.g., sulfonates) are attached to the first, second and fourth amino acid. For example, the peptide can be Ser-Ser-Y-Ser, wherein an anionic group is attached to the side chain of each serine residue and Y is any amino acid. In addition to peptides and peptide derivatives, single amino acids can be used as carriers in the A β 40 inhibitor of the invention. For example, cysteic acid, the sulfonate derivative of cysteine, can be used. Peptides such as disclosed in International Application No. WO 00/68263 may be used, also, e.g., Lys-Ile-Val-Phe-Phe-Ala (SEQ ID NO:1); Lys-Lys-Leu-Val-Phe-Phe-Ala (SEQ ID NO:2); Lys-Leu-Val-Phe-Phe-Ala (SEQ ID NO:3); Lys-Phe-Val-Phe-Phe-Ala (SEQ ID NO:4); Ala-Phe-Phe-Val-Leu-Lys (SEQ ID NO:5); Lys-Leu-Val-Phe (SEQ ID NO:6); Lys-Ala-Val-Phe-Phe-Ala (SEQ ID NO:7); Lys-Leu-Val-Phe-Phe (SEQ ID NO:8); Lys-Val-Val-Phe-Phe-Ala (SEQ ID NO:9); Lys-Ile-Val-Phe-Phe-Ala-NH₂ (SEQ ID NO:10); Lys-Leu-Val-Phe-Phe-Ala-NH₂ (SEQ ID NO:11); Lys-Phe-Val-Phe-Phe-Ala-NH₂ (SEQ ID NO:12); Ala-Phe-Phe-Val-Leu-Lys-NH₂ (SEQ ID NO:13); Lys-Leu-Val-Phe-NH₂ (SEQ ID NO:14); Lys-Ala-Val-Phe-Phe-Ala-NH₂ (SEQ ID NO:15); Lys-Leu-Val-Phe-Phe-NH₂ (SEQ ID NO:16); Lys-Val-Val-Phe-Phe-Ala-NH₂ (SEQ ID NO:17); Lys-Leu-Val-Phe-Phe-Ala-Gln (SEQ ID NO:18); Lys-Leu-Val-Phe-Phe-Ala-Gln-NH₂ (SEQ ID NO:19); His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-NH₂ (SEQ ID NO:20); Asp-Asp-Asp (SEQ ID NO:21); Lys-Val-Asp-Asp-Gln-Asp (SEQ ID NO:22); His-His-Gln-Lys (SEQ ID NO:23); and Gln-Lys-Leu-Val-Phe-Phe-NH₂ (SEQ ID NO:24).

The term "aliphatic group" is intended to include organic compounds characterized by straight or branched chains, typically having between 1 and 22 carbon atoms. Aliphatic groups include alkyl groups, alkenyl groups and alkynyl groups. In complex structures, the chains can be branched or cross-linked. Alkyl groups include saturated hydrocarbons having one or more carbon atoms, including straight-chain alkyl groups and branched-chain alkyl groups. Such hydrocarbon moieties may be substituted on one or more carbons with, for example, a halogen, a hydroxyl, a thiol, an amino, an alkoxy, an alkylcarboxy, an alkylthio, or a nitro group. Unless the number of carbons is otherwise specified, "lower aliphatic" as used herein means an aliphatic group, as defined above (e.g., lower alkyl, lower alkenyl, lower alkynyl), but having from one to six carbon atoms. Representatives of such lower aliphatic groups, e.g., lower alkyl groups, are methyl, ethyl, n-propyl, isopropyl, 2-chloropropyl, n-butyl, sec-butyl, 2-aminobutyl, isobutyl, tert-butyl, 3-thiopentyl, and the like. As used herein, the term "amino" means -NH₂; the term "nitro" means -NO₂; the term "halogen" designates -F, -Cl, -Br or -I; the term "thiol"

means SH; and the term "hydroxyl" means -OH. Thus, the term "alkylamino" as used herein means -NHR wherein R is an alkyl group as defined above. The term "alkylthio" refers to -SR, wherein R is an alkyl group as defined above. The term "alkylcarboxyl" as used herein means -COOR, wherein R is an alkyl group as defined above. The term "alkoxy" as used herein means -OR, wherein R is an alkyl group as defined above. Representative alkoxy groups include methoxy, ethoxy, propoxy, tert-butoxy and the like. The terms "alkenyl" and "alkynyl" refer to unsaturated aliphatic groups analogous to alkyls, but which contain at least one double or triple bond respectively.

The term "alicyclic group" is intended to include closed ring structures of three or more carbon atoms. Aicyclic groups include cycloparaffins or naphthenes which are saturated cyclic hydrocarbons, cycloolefins which are unsaturated with two or more double bonds, and cycloacetylenes which have a triple bond. They do not include aromatic groups. Examples of cycloparaffins include cyclopropane, cyclohexane, and cyclopentane. Examples of cycloolefins include cyclopentadiene and cyclooctatetraene. Aicyclic groups also include fused ring structures and substituted alicyclic groups such as alkyl substituted alicyclic groups. In the instance of the alicyclics such substituents can further comprise a lower alkyl, a lower alkenyl, a lower alkoxy, a lower alkylthio, a lower alkylamino, a lower alkylcarboxyl, a nitro, a hydroxyl, -CF₃, -CN, or the like.

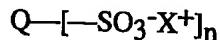
The term "heterocyclic group" is intended to include closed ring structures in which one or more of the atoms in the ring is an element other than carbon, for example, nitrogen, or oxygen. Heterocyclic groups can be saturated or unsaturated and heterocyclic groups such as pyrrole and furan can have aromatic character. They include fused ring structures such as quinoline and isoquinoline. Other examples of heterocyclic groups include pyridine and purine. Heterocyclic groups can also be substituted at one or more constituent atoms with, for example, a halogen, a lower alkyl, a lower alkenyl, a lower alkoxy, a lower alkylthio, a lower alkylamino, a lower alkylcarboxyl, a nitro, a hydroxyl, -CF₃, -CN, or the like.

The term "aromatic group" is intended to include unsaturated cyclic hydrocarbons containing one or more rings. Aromatic groups include 5- and 6-membered single-ring groups which may include from zero to four heteroatoms, for example, benzene, pyrrole, furan, thiophene, imidazole, oxazole, thiazole, triazole, pyrazole, pyridine, pyrazine, pyridazine and pyrimidine, and the like. The aromatic ring may be substituted at one or more ring positions

with, for example, a halogen, a lower alkyl, a lower alkenyl, a lower alkoxy, a lower alkylthio, a lower alkylamino, a lower alkylcarboxyl, a nitro, a hydroxyl, -CF₃, -CN, or the like.

In a preferred embodiment of the method of the invention, the A_β40 inhibitor administered to the subject is comprised of at least one sulfonate group covalently attached to a carrier group, or a pharmaceutically acceptable salt thereof. Accordingly, an A_β40 inhibitor can have the structure:

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wherein Q is a carrier group; X⁺ is a cationic group; and n is an integer. Suitable carrier groups and cationic groups are those described hereinbefore. The number of sulfonate groups 10 ("n") is selected such that the biodistribution of the compound for an intended target site is not prevented while maintaining activity of the compound as discussed earlier. In one embodiment, n is an integer between 1 and 10. In another embodiment, n is an integer between 3 and 8. As described earlier, an A_β40 inhibitor with multiple sulfonate groups can have the sulfonate groups spaced such that the compound interacts optimally with an HSPG binding site within the 15 A_β40 peptide.

In preferred embodiments, the carrier group for a sulfonate(s) is a lower aliphatic group (e.g., a lower alkyl, lower alkenyl or lower alkynyl), a heterocyclic group, and group derived from a disaccharide, a polymer or a peptide or peptide derivative. Furthermore, the carrier can be substituted, e.g., with one or more amino, nitro, halogeno, sulphydryl or hydroxyl groups. In 20 certain embodiments, the carrier for a sulfonate(s) is an aromatic group.

Examples of suitable sulfonated polymeric A_β40 inhibitors include poly(2-acrylamido-2-methyl-1-propanesulfonic acid); poly(2-acrylamido-2-methyl-1-propanesulfonic acid-co-acrylonitrile); poly(2-acrylamido-2-methyl-1-propanesulfonic acid-co-styrene); poly(vinylsulfonic acid); poly(4-styrenesulfonic acid); a sulfonic acid derivative of poly(acrylic acid); a sulfonic acid derivative of poly(methyl acrylate); a sulfonic acid derivative of poly(methyl methacrylate); and pharmaceutically acceptable salts thereof.

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A preferred sulfonated polymer is poly(vinylsulfonic acid) (PVS) or a pharmaceutically acceptable salt thereof, preferably the sodium salt thereof. In one embodiment, PVS having a molecular weight of about 800-1000 Daltons is used. PVS may be used as a mixture of isomers 30 or as a single active isomer.

Preferred sulfonated saccharides include 5-deoxy-1,2-O-isopropylidene- α -D-xylofuranose-5-sulfonic acid (XXIII, shown as the sodium salt).

Preferred lower aliphatic sulfonated A β 40 inhibitors for use in the invention include ethanesulfonic acid; 2-aminoethanesulfonic acid (taurine); cysteic acid (3-sulfoalanine or α -amino- β -sulfopropionic acid); 1-propanesulfonic acid; 1,2-ethanedisulfonic acid; 1,3-propanedisulfonic acid; 1,4-butanedisulfonic acid; 1,5-pentanedisulfonic acid; and 4-hydroxy-1-butanesulfonic acid (VIII, shown as the sodium salt); and pharmaceutically acceptable salts thereof. Other aliphatic sulfonated A β 40 inhibitors contemplated for use in the invention include 1-butanesulfonic acid (XLVII, shown as the sodium salt), 2-propanesulfonic acid (XLIX, shown as the sodium salt), 3-pentanesulfonic acid (L, shown as the sodium salt), 4-heptanesulfonic acid (LII, shown as the sodium salt), 1-decanesulfonic acid (XLVIII, shown as the sodium salt); and pharmaceutically acceptable salts thereof. Sulfonated substituted aliphatic A β 40 inhibitors contemplated for use in the invention include 3-amino-1-propanesulfonic acid (XXII, shown as the sodium salt), 3-hydroxy-1-propanesulfonic acid sulfate (XXXV, shown as the disodium salt), 1,7-dihydroxy-4-heptanesulfonic acid (LIII, shown as the sodium salt); and pharmaceutically acceptable salts thereof. Yet other sulfonated compounds contemplated for use in the invention include 2-[(4-pyridinyl)amido]ethanesulfonic acid (LIV, depicted as the sodium salt), and pharmaceutically acceptable salts thereof.

Preferred heterocyclic sulfonated A β 40 inhibitors include 3-(N-morpholino)-1-propanesulfonic acid; and tetrahydrothiophene-1,1-dioxide-3,4-disulfonic acid; and pharmaceutically acceptable salts thereof.

Aromatic sulfonated A β 40 inhibitors include 1,3-benzenedisulfonic acid (XXXVI, shown as the disodium salt), 2,5-dimethoxy-1,4-benzenedisulfonic acid (depicted as the disodium salt, XXXVII, or the dipotassium salt, XXXIX), 4-amino-3-hydroxy-1-naphthalenesulfonic acid (XLIII), 3,4-diamino-1-naphthalenesulfonic acid (XLIV); and pharmaceutically acceptable salts thereof.

In another embodiment of the method of the invention, the A β 40 inhibitor administered to the subject is comprised of at least one sulfate group covalently attached to a carrier group, or a pharmaceutically acceptable salt thereof. Accordingly, the A β 40 inhibitor can have the



wherein Q is a carrier group; X⁺ is a cationic group; and n is an integer. Suitable carriers and cationic groups are those described hereinbefore. The number of sulfate groups ("n") is selected such that the biodistribution of the compound for an intended target site is not prevented while maintaining activity of the Aβ40 inhibitor as discussed earlier. In one embodiment, n is an integer between 1 and 10. In another embodiment, n is an integer between 3 and 8. As described earlier, an Aβ40 inhibitor with multiple sulfate groups can have the sulfate groups spaced such that the compound interacts optimally with a GAG binding site within an Aβ peptide.

In preferred embodiments, the carrier group for a sulfate(s) is a lower aliphatic group (e.g., a lower alkyl, lower alkenyl or lower alkynyl), an aromatic group, a group derived from a disaccharide, a polymer or a peptide or peptide derivative. Furthermore, the carrier can be substituted, e.g., with one or more amino, nitro, halogeno, sulfhydryl or hydroxyl groups.

Examples of suitable sulfated polymeric Aβ40 inhibitors include poly(2-acrylamido-2-methyl-1-propyl sulfuric acid); poly(2-acrylamido-2-methyl-1-propyl sulfuric acid-co-acrylonitrile); poly(2-acrylamido-2-methyl-propyl sulfuric acid-co-styrene); poly(vinylsulfuric acid); poly(sodium 4-styrenesulfate); a sulfate derivative of poly(acrylic acid); a sulfate derivative of poly(methyl acrylate); a sulfate derivative of poly(methyl methacrylate); and a sulfate derivative of poly(vinyl alcohol); and pharmaceutically acceptable salts thereof.

A preferred sulfated polymer is poly(vinylsulfuric acid) or pharmaceutically acceptable salt thereof.

A preferred sulfated disaccharide is sucrose octasulfate or pharmaceutically acceptable salt thereof. Other sulfated saccharides contemplated for use in the invention include the acid form of methyl α-D-glucopyranoside 2,3-disulfate (XVI), methyl 4,6-O-benzylidene-α-D-glucopyranoside 2,3-disulfate (XVII), 2,3,4,3',4'-sucrose pentasulfate (XXXIII), 1,3:4,6-di-O-

disulfuric acid; and 1,4-butanediol monosulfuric acid; and pharmaceutically acceptable salts thereof. Other sulfated aliphatic A β 40 inhibitors contemplated for use in the invention include the acid form of 1,3-cyclohexanediol disulfate (XL), 1,3,5-heptanetriol trisulfate (XIX), 2-hydroxymethyl-1,3-propanediol trisulfate (XX), 2-hydroxymethyl-2-methyl-1,3-propanediol trisulfate (XXI), 1,3,5,7-heptanetetraol tetrasulfate (XLVI), 1,3,5,7,9-nonane pentasulfate (LI); and pharmaceutically acceptable salts thereof. Other sulfated A β 40 inhibitors contemplated for use in the invention include the acid form of 2-amino-2-hydroxymethyl-1,3-propanediol trisulfate (XXIV), 2-benzyloxy-1,3-propanediol disulfate (XXIX), 3-hydroxypropylsulfamic acid sulfate (XXX), 2,2'-iminoethanol disulfate (XXXI), N,N-bis(2-hydroxyethyl)sulfamic acid disulfate (XXXII); and pharmaceutically acceptable salts thereof.

Preferred heterocyclic sulfated A β 40 inhibitors include 3-(N-morpholino)-1-propyl sulfuric acid; and tetrahydrothiophene-3,4-diol-1,1-dioxide disulfuric acid; and pharmaceutically acceptable salts thereof.

The invention further contemplates the use of prodrugs which are converted *in vivo* to the A β 40 inhibitors used in the methods of the invention (see, e.g., R.B. Silverman, 1992, "The Organic Chemistry of Drug Design and Drug Action", Academic Press, Chp. 8). Such prodrugs can be used to alter the biodistribution (e.g., to allow compounds which would not typically cross the blood-brain barrier to cross the blood-brain barrier) or the pharmacokinetics of the A β 40 inhibitor. For example, an anionic group, e.g., a sulfate or sulfonate, can be esterified, e.g., with a methyl group or a phenyl group, to yield a sulfate or sulfonate ester. When the sulfate or sulfonate ester is administered to a subject, the ester is cleaved, enzymatically or non-enzymatically, reductively or hydrolytically, to reveal the anionic group. Such an ester can be cyclic, e.g., a cyclic sulfate or sultone, or two or more anionic moieties may be esterified through a linking group. Exemplary cyclic A β 40 inhibitors include, for example, 2-sulfobenzoic acid cyclic anhydride (LV), 1,3-propane sultone (LVI), 1,4-butane sultone (LVII), 1,3-butanediol cyclic sulfate (LVIII), α -chloro- α -hydroxy- α -toluenesulfonic acid γ -sultone (LIX), and 6-nitronaphth-[1,8-cd]-1,2-oxathiole 2,2-dioxide (LX). In a preferred embodiment, the prodrug is a cyclic sulfate or sultone. An anionic group can be esterified with moieties (e.g., acyloxymethyl esters) which are cleaved to reveal an intermediate A β 40 inhibitor which subsequently decomposes to yield the active A β 40 inhibitor. In another embodiment, the prodrug is a reduced form of a sulfate or sulfonate, e.g., a thiol, which is oxidized *in vivo* to the

A β 40 inhibitor. Furthermore, an anionic moiety can be esterified to a group which is actively transported *in vivo*, or which is selectively taken up by target organs. The ester can be selected to allow specific targeting of the A β 40 inhibitors to particular organs, as described below for carrier moieties.

5 Carrier groups useful in the A β 40 inhibitors include groups previously described; e.g., aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups, groups derived from carbohydrates, polymers, peptides, peptide derivatives, or combinations thereof. Suitable polymers include substituted and unsubstituted vinyl, acryl, styrene and carbohydrate-derived polymers and copolymers and salts thereof. Preferred carrier groups include a lower alkyl
10 group, a heterocyclic group, a group derived from a disaccharide, a polymer, a peptide, or peptide derivative.

Carrier groups useful in the present invention may also include moieties which allow the A β 40 inhibitor to be selectively delivered to a target organ or organs. For example, for a desirable delivery of an A β 40 inhibitor to the brain, the carrier group may include a moiety
15 capable of targeting the A β 40 inhibitor to the brain, by either active or passive transport (a "targeting moiety"). Illustratively, the carrier group may include a redox moiety, as described in, for example, U.S. Patents 4,540,564 and 5,389,623, both to Bodor. These patents disclose drugs linked to dihydropyridine moieties which can enter the brain, where they are oxidized to a charged pyridinium species which is trapped in the brain. Thus, drug accumulates in the brain.

20 Exemplary pyridine/dihydropyridine compounds of the invention include sodium 2-(nicotinylamido)-ethanesulfonate (LXII), and 1-(3-sulfopropyl)-pyridinium betaine (LXIII). Other carrier moieties include groups, such as those derived from amino acids or thyroxine, which can be passively or actively transported *in vivo*. An illustrative compound is phenylalanyltaurine (LXIX), in which a taurine molecule is conjugated to a phenylalanine (a
25 large neutral amino acid). Such a carrier moiety can be metabolically removed *in vivo*, or can remain intact as part of an active A β 40 inhibitor. Structural mimics of amino acids (and other actively transported moieties) are also useful in the invention (e.g., 1-(aminomethyl)-1-(sulfomethyl)-cyclohexane (LXX)). Other exemplary amino acid mimetics include p-(sulfomethyl)phenylalanine (LXXII), p-(1,3-disulfoprop-2-yl)phenylalanine (LXXIII), and O-
30 (1,3-disulfoprop-2-yl)tyrosine (LXXIV). Exemplary thyroxine mimetics include compounds LXV, LXVI, and LXVII. Many targeting moieties are known, and include, for example,

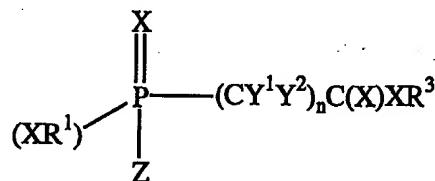
asialoglycoproteins (see, e.g., Wu, U.S. Patent No. 5,166,320) and other ligands which are transported into cells via receptor-mediated endocytosis (see below for further examples of targeting moieties which may be covalently or non-covalently bound to a carrier molecule). Furthermore, the A β 40 inhibitors of the invention may bind to amyloidogenic proteins, e.g., A β 40, in the circulation and thus be transported to the site of action.

The targeting and prodrug strategies described above can be combined to produce an A β 40 inhibitor that can be transported as a prodrug to a desired site of action and then unmasked to reveal an active A β 40 inhibitor. For example, the dihydropyridine strategy of Bodor (see *supra*) can be combined with a cyclic prodrug, as for example in the compound 2-10 (1-methyl-1,4-dihydronicotinoyl)amidomethyl-propanesultone (LXXI).

In one embodiment, the A β 40 inhibitor in the pharmaceutical compositions is a sulfonated polymer, for example poly(2-acrylamido-2-methyl-1-propanesulfonic acid); poly(2-acrylamido-2-methyl-1-propanesulfonic acid-co-acrylonitrile); poly(2-acrylamido-2-methyl-1-propanesulfonic acid-co-styrene); poly(vinylsulfonic acid); poly(4-styrenesulfonic acid); a sulfonate derivative of poly(acrylic acid); a sulfonate derivative of poly(methyl acrylate); a sulfonate derivative of poly(methyl methacrylate); and a sulfonate derivative of poly(vinyl alcohol); and pharmaceutically acceptable salts thereof.

In another embodiment, the A β 40 inhibitor in the pharmaceutical compositions is a sulfated polymer, for example poly(2-acrylamido-2-methyl-1-propyl sulfuric acid); poly(2-acrylamido-2-methyl-1-propyl sulfuric acid-co-acrylonitrile); poly(2-acrylamido-2-methyl-1-propyl sulfuric acid-co-styrene); poly(vinyl sulfuric acid); poly(4-styrenesulfate); a sulfate derivative of poly(acrylic acid); a sulfate derivative of poly(methyl acrylate); a sulfate derivative of poly(methyl methacrylate); and pharmaceutically acceptable salts thereof.

The A β 40 inhibitor can also have the structure:



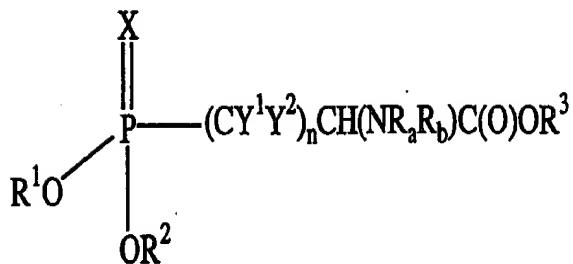
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wherein Z is XR² or R⁴, R¹ and R² are each independently hydrogen, a substituted or unsubstituted aliphatic group (preferably a branched or straight-chain aliphatic moiety having

from 1 to 24 carbon atoms in the chain; or an unsubstituted or substituted cyclic aliphatic moiety having from 4 to 7 carbon atoms in the aliphatic ring; preferred aliphatic and cyclic aliphatic groups are alkyl groups, more preferably lower alkyl), an aryl group, a heterocyclic group, or a salt-forming cation; R³ is hydrogen, lower alkyl, aryl, or a salt-forming cation; X is, independently for each occurrence, O or S; R⁴ is hydrogen, lower alkyl, aryl or amino; Y¹ and Y² are each independently hydrogen, halogen (e.g., F, Cl, Br, or I), lower alkyl, amino (including alkylamino, dialkylamino, arylamino, diarylamino, and alkylarylamino), hydroxy, alkoxy, or aryloxy; and n is an integer from 0 to 12 (more preferably 0 to 6, more preferably 0 or 1). These compounds are described in U.S. Patent No. 5,869,469, the contents of which is incorporated herein by reference.

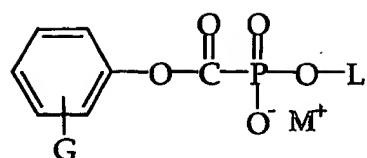
Preferred A β 40 inhibitors for use in the invention include compounds in which both R¹ and R² are pharmaceutically acceptable salt-forming cations. It will be appreciated that the stoichiometry of an anionic compound to a salt-forming counterion (if any) will vary depending on the charge of the anionic portion of the compound (if any) and the charge of the counterion.

In a particularly preferred embodiment, R¹, R² and R³ are each independently a sodium, potassium or calcium cation. In certain embodiments in which at least one of R¹ and R² is an aliphatic group, the aliphatic group has between 1 and 10 carbons atoms in the straight or branched chain, and is more preferably a lower alkyl group. In other embodiments in which at least one of R¹ and R² is an aliphatic group, the aliphatic group has between 10 and 24 carbons atoms in the straight or branched chain. In certain preferred embodiments, n is 0 or 1; more



wherein R^1 , R^2 , R^3 , Y^1 , Y^2 , and X are as defined above, R_a and R_b are each independently hydrogen, alkyl, aryl, or heterocyclyl, or R_a and R_b , taken together with the nitrogen atom to which they are attached, form a cyclic moiety having from 3 to 8 atoms in the ring, and n is an integer from 0 to 6. In certain preferred embodiments, R_a and R_b are each hydrogen. In certain preferred embodiments, a compound of the invention comprises an α -amino acid (or α -amino acid ester), more preferably a L- α -amino acid or ester.

The Z , R^1 , R^2 , R^3 , Y^1 , Y^2 and X groups are each independently selected such that the biodistribution of the A β 40 inhibitor for an intended target site is not prevented while maintaining activity of the A β 40 inhibitor. For example, the number of anionic groups (and the overall charge on the therapeutic compound) should not be so great as to prevent traversal of an anatomical barrier, such as a cell membrane, or entry across a physiological barrier, such as the blood-brain barrier, in situations where such properties are desired. For example, it has been reported that esters of phosphonoformate have biodistribution properties different from, and in some cases superior to, the biodistribution properties of phosphonoformate (see, e.g., U.S. Patent Nos. 4,386,081 and 4,591,583 to Helgstrand et al., and U.S. Patent Nos. 5,194,654 and 5,463,092 to Hostetler et al.). Thus, in certain embodiments, at least one of R^1 and R^2 is an aliphatic group (more preferably an alkyl group), in which the aliphatic group has between 10 and 24 carbons atoms in the straight or branched chain. The number, length, and degree of branching of the aliphatic chains can be selected to provide a desired characteristic, e.g., lipophilicity. In other embodiments, at least one of R^1 and R^2 is an aliphatic group (more preferably an alkyl group), in which the aliphatic group has between 1 and 10 carbons atoms in



wherein G represents hydrogen or one or more substituents on the aryl ring (e.g., alkyl, aryl, halogen, amino, and the like) and L is a substituted alkyl group (in certain embodiments, preferably a lower alkyl), more preferably a hydroxy-substituted alkyl or an alkyl substituted with a nucleoside base, and M⁺ is a counter ion. In certain embodiments, G is hydrogen or an electron-donating group. In embodiments in which G is an electron-withdrawing group, G is preferably an electron withdrawing group at the meta position. The term "electron-withdrawing group" is known in the art, and, as used herein, refers to a group which has a greater electron-withdrawing than hydrogen. A variety of electron-withdrawing groups are known, and include halogens (e.g., fluoro, chloro, bromo, and iodo groups), nitro, cyano, and the like. Similarly, the term "electron-donating group", as used herein, refers to a group which is less electron-withdrawing than hydrogen. In embodiments in which G is an electron donating group, G can be in the ortho, meta or para position. In certain embodiments, M⁺ is a cationic species selected from, e.g., H⁺ and pharmaceutically acceptable organic or inorganic ions, including, without limitation, Na⁺, K⁺, NH₄⁺, Ca⁺², RNH₃⁺, RR'NH₂⁺. In one preferred embodiment, M⁺ is an anilinium ion.

In certain preferred embodiments, L may be one of the following moieties :

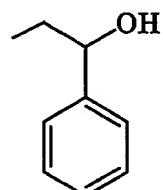
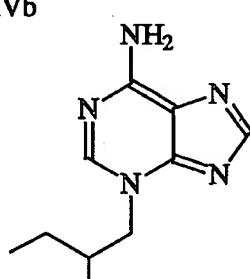
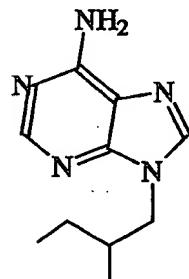
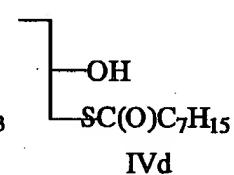
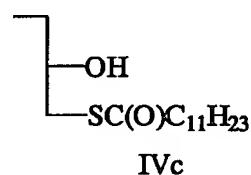
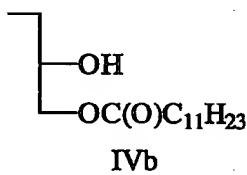
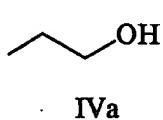


Table 1 lists data pertinent to the characterization of these compounds using art-
recognized techniques. The compounds IVa-IVg in Table 1 correspond to the following
structure, wherein L is a group selected from the above-listed (Groups IVa-IVg) having the
same number.

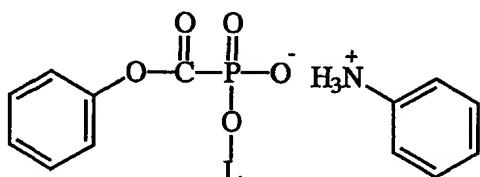


Table 1

<u>COMPOUND</u>	<u>^{31}P NMR</u>	<u>^{13}C NMR</u>	<u>FAB-MS(-)</u>
IVa	-6.33(DMSO-d ₆)	60.97 CH ₂ OH(d, J=6Hz) 66.76 CHOH(d, J=7.8Hz) 121.65, 121.78, 121.99, 125.71, 129.48, 129.57, 126.43 Aromatic CH 134.38 Aniline C-N 150.39 Phenyl C-O(d, J=7Hz) 171.57 P-C=O(d, J=234Hz)	245.2
IVb	-6.41(DMSO-d ₆)	13.94 CH ₃ 22.11, 24.40, 28.56, 28.72, 28.99, 29.00, 31.30, 33.43, -(CH ₂) ₁₀ ⁻ 65.03 CH ₂ -OC(O) 66.60 CH ₂ -OP(d, J=5.6Hz) 67.71 CH ₂ -OH(d, J=6 Hz) 121.73, 121.10, 125.64, 126.57, 129.40, 129.95, Aromatic CH 134.04 Aniline C-N 150.31 Phenyl C-O 171.44 P-C=O(d, J=6.7 Hz) 172.83 O-C=O	456
IVc	-6.46(DMSO-d ₆)	13.94 CH ₃ 22.11, 25.10, 28.68, 28.72, 28.85, 29.00, 30.76, 31.31, 32.10, -(CH ₂) ₁₀ ⁻ 43.36 CH ₂ -S 68.43 CH ₂ -OH 68.43 CH-OH(d, J=6.3 Hz) 68.76 P-O-CH ₂ -9d, J=5.8 Hz) 121.75, 122.03, 125.62, 126.37, 129.30, 129.53, Aromatic CH	471

<u>COMPOUND</u>	<u>³¹P NMR</u>	<u>¹³C NMR</u>	<u>FAB-MS(-)</u>	
IVd	-6.61(DMSO-d ₆)	13.94 CH ₃ 22.06, 25.14, 28.24, 28.35, 31.09, 32.14 -CH ₂) ₆ - 43.40 CH ₂ -S 68.50 P-O-CH ₂ -(d, J=5.8 Hz) 68.77 CH-OH(d, 6.4 Hz) 121.78, 122.59, 125.69, 127.06, 129.43, 129.59 Aromatic CH 133.39 Aniline C-N 150.38 Phenyl C-O(d, J=6.7 Hz) 171.47 P-C=O(d, J=234.4 Hz) 198.54 S-C=O		416
IVe	-5.76(D ₂ O)	N/A	N/A	
IVf	-7.00(DMSO-d ₆)	N/A	N/A	
IVg	-6.60(DMSO-D ₆)	70.84 CH ₂ -OH 72.17 CH-OH 121.68, 121.79, 121.85, 125.71 127.10, 127.92, 129.36, 129.50, 129.59 Aromatic CH 134.51 Aniline C-N 142.34 Aromatic C-CH 150.37 Phenyl C-O(d, J=6.2 Hz) 171.59 P-C=O(d, J=232.6 Hz)	321	

Note that the structure of some of the A_β40 inhibitors of this invention includes asymmetric carbon atoms. It is to be understood accordingly that the isomers (e.g., enantiomers and diastereomers) arising from such asymmetry are included within the scope of this

5 invention. Such isomers can be obtained in substantially pure form by classical separation techniques and by sterically controlled synthesis. For the purposes of this application, unless expressly noted to the contrary, an A_β40 inhibitor shall be construed to include both the R or S stereoisomers at each chiral center.

In certain embodiments, an A_β40 inhibitor of the invention comprises a cation (i.e., in
10 certain embodiments, at least one of R¹, R² or R³ is a cation). If the cationic group is hydrogen, H⁺, then the A_β40 inhibitor is considered an acid, e.g., phosphonoformic acid. If hydrogen is

replaced by a metal ion or its equivalent, the A β 40 inhibitor is a salt of the acid.

Pharmaceutically acceptable salts of the A β 40 inhibitor are within the scope of the invention.

For example, at least one of R¹, R² or R³ can be a pharmaceutically acceptable alkali metal

(e.g., Li, Na, or K), ammonium cation, alkaline earth cation (e.g., Ca²⁺, Ba²⁺, Mg²⁺), higher

5 valency cation, or polycationic counter ion (e.g., a polyammonium cation). (See, e.g., Berge et al. (1977) "Pharmaceutical Salts", *J. Pharm. Sci.* 66:1-19). It will be appreciated that the stoichiometry of an anionic compound to a salt-forming counterion (if any) will vary depending on the charge of the anionic portion of the compound (if any) and the charge of the counterion.

Preferred pharmaceutically acceptable salts include a sodium, potassium or calcium salt, but

10 other salts are also contemplated within their pharmaceutically acceptable range.

The term "pharmaceutically acceptable esters" refers to the relatively non-toxic, esterified products of the A β 40 inhibitors of the present invention. These esters can be prepared *in situ* during the final isolation and purification of the A β 40 inhibitors or by separately reacting the purified A β 40 inhibitor in its free acid form or hydroxyl with a suitable esterifying agent; either of which are methods known to those skilled in the art. Carboxylic acids and phosphonic acids can be converted into esters according to methods well known to one of ordinary skill in the art, e.g., via treatment with an alcohol in the presence of a catalyst. A preferred ester group (e.g., when R³ is lower alkyl) is an ethyl ester group.

The term "alkyl" refers to the saturated aliphatic groups, including straight-chain alkyl groups, branched-chain alkyl groups, cycloalkyl (alicyclic) groups, alkyl substituted cycloalkyl groups, and cycloalkyl substituted alkyl groups. In preferred embodiments, a straight chain or branched chain alkyl has 30 or fewer carbon atoms in its backbone (e.g., C₁-C₃₀ for straight chain, C₃-C₃₀ for branched chain), and more preferably 20 or fewer. Likewise, preferred cycloalkyls have from 4-10 carbon atoms in their ring structure, and more preferably have 4-7 carbon atoms in the ring structure. The term "lower alkyl" refers to alkyl groups having from 1

aryloxycarbonyloxy, carboxylate, alkylcarbonyl, alkoxycarbonyl, aminocarbonyl, alkylthiocarbonyl, alkoxy, phosphate, phosphonato, phosphinato, cyano, amino (including alkyl amino, dialkylamino, arylamino, diarylamino, and alkylarylamino), acylamino (including alkylcarbonylamino, arylcarbonylamino, carbamoyl and ureido), amidino, imino, sulfhydryl, 5 alkylthio, arylthio, thiocarboxylate, sulfate, sulfonato, sulfamoyl, sulfonamido, nitro, trifluoromethyl, cyano, azido, heterocyclyl, aralkyl, or an aromatic or heteroaromatic moiety. It will be understood by those skilled in the art that the moieties substituted on the hydrocarbon chain can themselves be substituted, if appropriate. Cycloalkyls can be further substituted, e.g., with the substituents described above. An "aralkyl" moiety is an alkyl substituted with an aryl 10 (e.g., phenylmethyl (benzyl)).

The term "alkoxy", as used herein, refers to a moiety having the structure -O-alkyl, in which the alkyl moiety is described above.

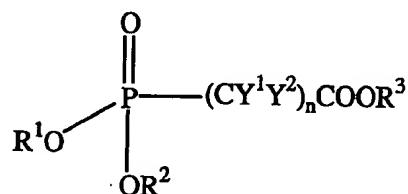
The term "aryl" as used herein includes 5- and 6-membered single-ring aromatic groups that may include from zero to four heteroatoms, for example, unsubstituted or substituted 15 benzene, pyrrole, furan, thiophene, imidazole, oxazole, thiazole, triazole, pyrazole, pyridine, pyrazine, pyridazine and pyrimidine, and the like. Aryl groups also include polycyclic fused aromatic groups such as naphthyl, quinolyl, indolyl, and the like. The aromatic ring can be substituted at one or more ring positions with such substituents, e.g., as described above for alkyl groups. Preferred aryl groups include unsubstituted and substituted phenyl groups.

20 The term "aryloxy", as used herein, refers to a group having the structure -O-aryl, in which the aryl moiety is as defined above.

The term "amino," as used herein, refers to an unsubstituted or substituted moiety of the formula -NR_aR_b, wherein R_a and R_b are each independently hydrogen, alkyl, aryl, or heterocyclyl, or R_a and R_b taken together with the nitrogen atom to which they are attached

from 10 to 24 carbons in the aliphatic chain, e.g., the long-chain aliphatic moiety is an aliphatic chain of a fatty acid (preferably a naturally-occurring fatty acid). Representative long-chain aliphatic moieties include the aliphatic chains of stearic acid, oleic acid, linolenic acid, and the like.

5 In certain embodiments, the A_β40 inhibitor of the invention can have the structure:



wherein R¹ and R² are each independently hydrogen, an aliphatic group (preferably a branched or straight-chain aliphatic moiety having from 1 to 24 carbon atoms, more preferably 10-24 carbon atoms, in the chain; or an unsubstituted or substituted cyclic aliphatic moiety having from 4 to 7 carbon atoms in the aliphatic ring), an aryl group, a heterocyclic group, or a salt-forming cation; R³ is hydrogen, lower alkyl, aryl, or a salt-forming cation; Y¹ and Y² are each independently hydrogen, halogen (e.g., F, Cl, Br, or I), lower alkyl, hydroxy, alkoxy, or aryloxy; and n is an integer from 0 to 12. Preferred A_β40 inhibitors for use in the invention include compounds wherein both R¹ and R² are pharmaceutically acceptable salt-forming cations. In a particularly preferred embodiment, R¹, R² and R³ are each independently a sodium, potassium or calcium cation, and n is 0. In certain preferred embodiments of the therapeutic compounds, Y¹ and Y² are each hydrogen. Particularly preferred A_β40 inhibitors are salts of phosphonoformate. Trisodium phosphonoformate (foscarnet sodium or Foscavir[®]) is commercially available (e.g., from Astra), and its clinical pharmacology has been investigated (see, e.g., "Physician's Desk Reference", 51st Ed., pp. 541-545 (1997)).

In another embodiment, the A_β40 inhibitor used in the invention can be an aminophosphonate, a bisphosphonate, a phosphonocarboxylate derivative, a phosphonate derivative, or a phosphono carbohydrate. For example, the A_β40 inhibitor can be one of the compounds described in Tables III and IV.

25 **Pharmaceutically Acceptable Formulations**

In the methods of the invention, the A_β40 inhibitor can be administered in a pharmaceutically acceptable formulation. The present invention pertains to any

pharmaceutically acceptable formulations, such as synthetic or natural polymers in the form of macromolecular complexes, nanocapsules, microspheres, or beads, and lipid-based formulations including oil-in-water emulsions, micelles, mixed micelles, synthetic membrane vesicles, and resealed erythrocytes.

5 In one embodiment, the pharmaceutically acceptable formulations comprise a polymeric matrix.

The terms "polymer" or "polymeric" are art-recognized and include a structural framework comprised of repeating monomer units which is capable of delivering an A β 40 inhibitor, such that treatment of a targeted condition occurs. The terms also include co-
10 polymers and homopolymers e.g., synthetic or naturally occurring. Linear polymers, branched polymers, and cross-linked polymers are also meant to be included.

For example, polymeric materials suitable for forming the pharmaceutically acceptable formulation employed in the present invention, include naturally derived polymers such as albumin, alginate, cellulose derivatives, collagen, fibrin, gelatin, and polysaccharides, as well as
15 synthetic polymers such as polyesters (PLA, PLGA), polyethylene glycol, poloxomers, polyanhydrides, and pluronic. These polymers are biocompatible and biodegradable without producing any toxic byproducts of degradation, and they possess the ability to modify the manner and duration of A β 40 inhibitor release by manipulating the polymer's kinetic characteristics. As used herein, the term "biodegradable" means that the polymer will degrade
20 over time by the action of enzymes, by hydrolytic action and/or by other similar mechanisms in the body of the subject. As used herein, the term "biocompatible" means that the polymer is compatible with a living tissue or a living organism by not being toxic or injurious and by not causing an immunological rejection.

Polymers can be prepared using methods known in the art (Sandler, S. R.; Karo, W.
25 *Polymer Syntheses*; Harcourt Brace: Boston, 1994; Shalaby, W.; Ikada, Y.; Langer, R.; Williams, J. *Polymers of Biological and Biomedical Significance (ACS Symposium Series 540)*; American Chemical Society: Washington, DC, 1994). Polymers can be designed to be flexible; the distance between the bioactive side-chains and the length of a linker between the polymer backbone and the group can be controlled. Other suitable polymers and methods for their
30 preparation are described in U.S. Patent Nos. 5,455,044 and 5,576,018.

The polymeric formulations are preferably formed by dispersion of the A β 40 inhibitor within liquefied polymer, as described in U.S. Patent No. 4,883,666, the teachings of which are incorporated herein by reference, or by such methods as bulk polymerization, interfacial polymerization, solution polymerization and ring polymerization as described in Odian G., 5 *Principles Of Polymerization And Ring Opening Polymerization*, 2nd ed., John Wiley & Sons, New York, 1981. The properties and characteristics of the formulations are controlled by varying such parameters as the reaction temperature, concentrations of polymer and A β 40 inhibitor, types of solvent used, and reaction times.

In addition to the A β 40 inhibitor and the pharmaceutically acceptable polymer, the 10 pharmaceutically acceptable formulation used in the method of the invention can comprise additional pharmaceutically acceptable carriers and/or excipients. As used herein, “pharmaceutically acceptable carrier” includes any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents, and the like that are physiologically compatible. For example, the carrier can be suitable for injection into the 15 cerebrospinal fluid. Excipients include pharmaceutically acceptable stabilizers and disintegrants.

The A β 40 inhibitor can be encapsulated in one or more pharmaceutically acceptable polymers, to form a microcapsule, microsphere, or microparticle, terms used herein interchangeably. Microcapsules, microspheres, and microparticles are conventionally free-flowing powders consisting of spherical particles of 2 mm or less in diameter, usually 500 μ m or less in diameter. Particles less than 1 μ m are conventionally referred to as nanocapsules, 20 nanoparticles or nanospheres. For the most part, the difference between a microcapsule and a nanocapsule, a microsphere and a nanosphere, or microparticle and nanoparticle is size; generally there is little, if any, difference between the internal structure of the two. In one 25 aspect of the present invention, the mean average diameter is less than about 45 μ m, preferably less than 20 μ m, and more preferably between about 0.1 and 10 μ m.

In another embodiment, the pharmaceutically acceptable formulations comprise lipid-based formulations. Any of the known lipid-based drug delivery systems can be used in the practice of the invention. For instance, multivesicular liposomes (MVL), multilamellar 30 liposomes (also known as multilamellar vesicles or “MLV”), unilamellar liposomes, including small unilamellar liposomes (SULV), and large unilamellar liposomes (“LULV”).

unilamellar liposomes (also known as large unilamellar vesicles or "LUV"), can all be used so long as a sustained release rate of the encapsulated A β 40 inhibitor can be established. In one embodiment, the lipid-based formulation can be a multivesicular liposome system. Methods of making controlled release multivesicular liposome drug delivery systems is described in PCT
5 Application Nos. US96/11642, US94/12957 and US94/04490.

The composition of the synthetic membrane vesicle is usually a combination of phospholipids, usually in combination with steroids, especially cholesterol. Other phospholipids or other lipids may also be used.

Examples of lipids useful in synthetic membrane vesicle production include
10 phosphatidylglycerols, phosphatidylcholines, phosphatidylserines, phosphatidylethanolamines, sphingolipids, cerebrosides, and gangliosides. Preferably phospholipids including egg phosphatidylcholine, dipalmitoylphosphatidylcholine, distearoylphosphatidylcholine, dioleoylphosphatidylcholine, dipalmitoylphosphatidylglycerol, and dioleoylphosphatidylglycerol are used.

15 In preparing lipid-based vesicles containing an A β 40 inhibitor, such variables as the efficiency of A β 40 inhibitor encapsulation, lability of the A β 40 inhibitor, homogeneity and size of the resulting population of vesicles, A β 40 inhibitor-to-lipid ratio, permeability, instability of the preparation, and pharmaceutical acceptability of the formulation should be considered (see Szoka, et al., *Annual Reviews of Biophysics and Bioengineering*, 9:467, 1980; Deamer, et al., in
20 *Liposomes*, Marcel Dekker, New York, 1983, 27; and Hope, et al., *Chem. Phys. Lipids*, 40:89, 1986, the contents of which are incorporated herein by reference).

Administration f the Pharmaceutically Acceptable Formulation

The A β 40 inhibitor may be administered to a subject, e.g., parenterally, e.g., intravenously, intradermally, subcutaneously, orally (e.g., via inhalation), transdermally
25 (topically), transmucosally, or rectally. In one embodiment, the A β 40 inhibitor is administered by introduction into the central nervous system of the subject, e.g., into the cerebrospinal fluid

Prior to introduction, the formulations can be sterilized with, preferably, gamma radiation or electron beam sterilization.

In another embodiment of the invention, the A_β40 inhibitor formulation is administered into a subject intrathecally. As used herein, the term "intrathecal administration" is intended to include delivering an A_β40 inhibitor formulation directly into the cerebrospinal fluid of a subject, by techniques including lateral cerebroventricular injection through a burrhole or cisternal or lumbar puncture or the like (described in Lazorthes et al. *Advances in Drug Delivery Systems and Applications in Neurosurgery*, 143-192 and Omaya et al., *Cancer Drug Delivery*, 1: 169-179, the contents of which are incorporated herein by reference). The term "lumbar region" is intended to include the area between the third and fourth lumbar (lower back) vertebrae. The term "cisterna magna" is intended to include the area where the skull ends and the spinal cord begins at the back of the head. The term "cerebral ventricle" is intended to include the cavities in the brain that are continuous with the central canal of the spinal cord. Administration of an A_β40 inhibitor to any of the above mentioned sites can be achieved by direct injection of the A_β40 inhibitor formulation or by the use of infusion pumps. For injection, the A_β40 inhibitor formulation of the invention can be formulated in liquid solutions, preferably in physiologically compatible buffers such as Hank's solution or Ringer's solution. In addition, the A_β40 inhibitor formulation may be formulated in solid form and re-dissolved or suspended immediately prior to use. Lyophilized forms are also included. The injection can be, for example, in the form of a bolus injection or continuous infusion (e.g., using infusion pumps) of the formulation.

Duration and Levels of Administration

In another embodiment of the method of the invention, the pharmaceutically acceptable formulation provides sustained delivery, e.g., "slow release" of the A_β40 inhibitor to a subject

cerebral amyloid angiopathy over time). Alternatively, sustained delivery of the A β 40 inhibitor may be demonstrated by detecting the presence of the A β 40 inhibitor *in vivo* over time.

In one embodiment, the pharmaceutically acceptable formulation provides sustained delivery of the A β 40 inhibitor to a subject for less than 30 days after the A β 40 inhibitor is administered to the subject. For example, the pharmaceutically acceptable formulation, e.g., "slow release" formulation, can provide sustained delivery of the A β 40 inhibitor to a subject for one, two, three or four weeks after the A β 40 inhibitor is administered to the subject. Alternatively, the pharmaceutically acceptable formulation may provide sustained delivery of the A β 40 inhibitor to a subject for more than 30 days after the A β 40 inhibitor is administered to the subject.

The pharmaceutical formulation, used in the method of the invention, contains a therapeutically effective amount of the A β 40 inhibitor. A "therapeutically effective amount" refers to an amount effective, at dosages and for periods of time necessary, to achieve the desired result. A therapeutically effective amount of the A β 40 inhibitor may vary according to factors such as the disease state, age, and weight of the subject, and the ability of the A β 40 inhibitor (alone or in combination with one or more other agents) to elicit a desired response in the subject. Dosage regimens may be adjusted to provide the optimum therapeutic response. A therapeutically effective amount is also one in which any toxic or detrimental effects of the A β 40 inhibitor are outweighed by the therapeutically beneficial effects. A non-limiting range for a therapeutically effective concentration of an A β 40 inhibitor is 100 μ M to 1 mM. It is to be further understood that for any particular subject, specific dosage regimens should be adjusted over time according to the individual need and the professional judgment of the person administering or supervising the administration of the A β 40 inhibitor and that dosage ranges set forth herein are exemplary only and are not intended to limit the scope or practice of the claimed invention.

In Vitro Treatment of Blood vessel wall cells

Blood vessel wall cells, or isolated blood vessel wall cells, can further be contacted with a therapeutically effective amount of a A β 40 inhibitor, *in vitro*. Accordingly, such cells can be isolated from a subject and grown *in vitro*, using techniques well known in the art. Briefly, a smooth muscle cell culture can be obtained by allowing smooth muscle cells to migrate out of

the tissue, e.g., mechanically or enzymatically, to produce a suspension of cells. For example, the enzymes trypsin, collagenase, elastase, hyaluronidase, DNase, pronase, dispase, or various combinations thereof can be used. Trypsin and pronase give the most complete disaggregation but may damage the cells. Collagenase and dispase give a less complete disaggregation but are 5 less harmful. Methods for isolating tissue such as neuronal tissue, and the disaggregation of tissue to obtain cells such as neuronal cells are described in Freshney R. I., *Culture of Animal Cells, A Manual of Basic Technique, Third Edition*, 1994, the contents of which are incorporated herein by reference.

Such cells can be subsequently contacted with an A β 40 inhibitor at levels and for a 10 duration of time as described above. Once inhibition of cerebral amyloid angiopathy has been achieved, these neuronal cells can be re-administered to the subject, e.g., by implantation.

States Characterized by CAA

The present invention further pertains to a method of treating a disease state characterized by cerebral amyloid angiopathy in a subject. As used herein, the term "state" is 15 art recognized and includes a disorder, disease or condition characterized by cerebral amyloid angiopathy. Examples of such disorders include Alzheimer's Disease, HCHWA-D, and hemorrhagic stroke.

The invention is further illustrated by the following examples, which should not be construed as further limiting. The contents of all references, patents and published patent 20 applications cited throughout this application are hereby incorporated by reference.

EXAMPLE 1

A compound of the invention is administered in a therapeutic amount to a subject having a clinical diagnosis of 'probable CAA', defined for present purposes as: multiple hemorrhages confined to the lobar brain regions diagnosed by CT or MRI scan and no other 25 cause of hemorrhage. The ability of the compound of the invention to prevent recurrence of CAA-related hemorrhages is determined by clinical exams (new neurologic symptoms or death with acute hemorrhage confirmed by CT scan or autopsy) or by gradient-echo MRI scans which mark the progression of CAA by the appearance of new hemorrhages. The ability of the compound to inhibit the progression of CAA can also be assessed through cognitive decline 30 (MMSE) or functional decline (NIHSS, FIM). The APOE-2 and APOE-4 are associated with

increasing risk and earlier age of first hemorrhage, but are neither specific nor sensitive for CAA.

EXAMPLE 2

The ability of compounds of the invention to inhibit CAA was measured in the following example. Nine week old hAPP transgenic mice were treated for a period of 8 weeks with two different concentrations of a compound of the present invention, 3-amino-1-propanesulfonic acid, sodium salt, 100 and 30 mg/kg. Mice were administered the compound for 8 weeks, after which they were sacrificed and their brains were perfused and processed for histological staining with Thioflavin S. This method may also be used as a screening method for determining activity of a candidate compound for inhibiting CAA.

The extent of CAA in brain sections obtained from these animals was qualitatively determined following staining. The extent of CAA, if any, was graded as follows:

- + Slight deposition
- ++ Moderate deposition
- +++ Severe deposition

The results shown in Table II, below, indicate that the test compound was effective in 1) reducing the number of mice showing CAA, and 2) showing an effect on the severity of the deposition seen in the brain vasculature of these animals.

Table II

Treatment	# animals in study	# of animals with CAA	CAA animals/total animals	CAA Severity		
				+	++	+++
Vehicle	16	15	15/16	5/15	9/15	1/15
30 mg/kg	11	10	10/11	6/10	4/10	—
100 mg/kg	15	10	10/15	9/10	—	1/10

Table III

Phosphonoacetic acid	
Phosphonoformic acid, trisodium salt hexahydrate	
Diethylphosphonoacetic acid	
3-[2-(1,2,3,4-Tetrahydroisoquinolinyl)]-1-propanephosphonic acid	
3-Aminopropylphosphonic acid	NH ₂ CH ₂ CH ₂ CH ₂ PO ₃ H ₂
Propylphosphonic acid	CH ₃ CH ₂ CH ₂ PO ₃ H ₂
Ethylphosphonic acid	CH ₃ CH ₂ PO ₃ H ₂
Methylphosphonic acid	CH ₃ PO ₃ H ₂
<i>tert</i> -Butylphosphonic acid	(CH ₃) ₃ CPO ₃ H ₂
Phenylphosphonic acid	
(<i>dl</i>)-2-Amino-3-phosphonopropanoic acid	
(1-Aminopropyl)phosphonic acid	
(<i>dl</i>)-2-Amino-5-phosphonopentanoic acid	
Diethyl phosphoramidate	
(<i>S</i>)-2-Amino-2-methyl-4-phosphonobutanoic acid	
D-(-)-2-Amino-4-phosphonobutanoic acid	

Table III

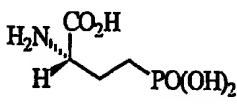
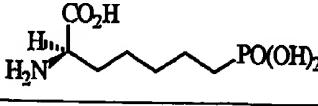
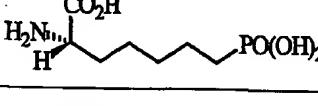
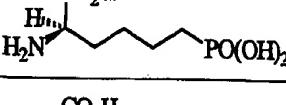
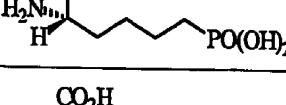
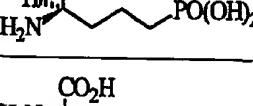
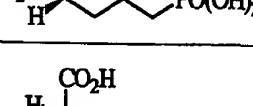
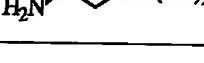
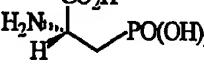
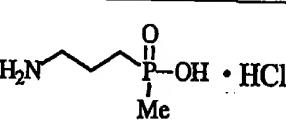
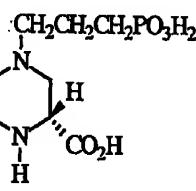
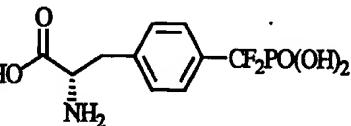
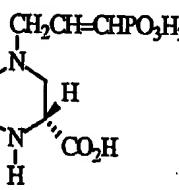
L-(+)-2-Amino-4-phosphonobutanoic acid	
D-(+)-2-Amino-7-phosphonoheptanoic acid	
L-(+)-2-Amino-7-phosphonoheptanoic acid	
D-(+)-2-Amino-6-phosphonohexanoic acid	
L-(+)-2-Amino-6-phosphonohexanoic acid	
D-(+)-2-Amino-4-phosphonopentanoic acid	
L-(+)-2-Amino-4-phosphonopentanoic acid	
D-(+)-2-Amino-3-phosphonopropanoic acid	
L-(+)-2-Amino-3-phosphonopropanoic acid	
3-Aminopropyl(methyl)phosphinic acid, hydrochloride	
(R)-(+)3-(2-Carboxypiperazin-4-yl)-propyl-1-phosphonic acid (D-CPP)	
L-4-[Difluoro(phosphono)methyl]-phenylalanine	
(R,E)-4-(3-Phosphonoprop-2-enyl)piperazine-2-carboxylic acid	

Table III

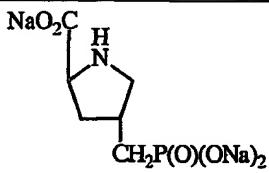
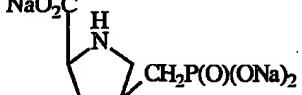
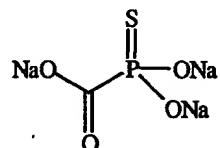
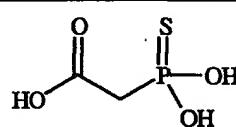
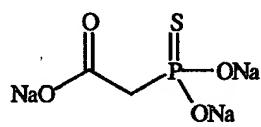
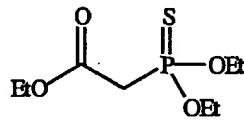
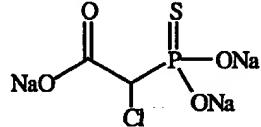
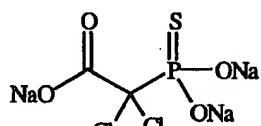
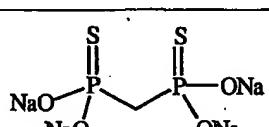
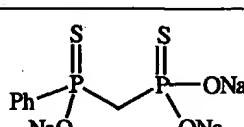
<i>trans</i> -L-4-Phosphonomethylproline, trisodium salt	
<i>cis</i> -L-4-Phosphonomethylproline, trisodium salt	
Thiophosphonoformic acid, trisodium salt	
Thiophosphonoacetic acid	
Thiophosphonoacetic acid, trisodium salt	
Thiophosphonoacetic acid, triethyl ester	
Chloro(thiophosphono)acetic acid, trisodium salt	
Dichloro(thiophosphono)acetic acid, trisodium salt	
Thiophosphonomethylthiophosphonic acid, tetrasodium salt	
Phenylthiophosphinomethylthio-phosphonic acid, trisodium salt	
Propylthiophosphonic acid	

Table III

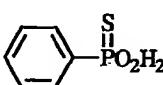
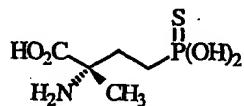
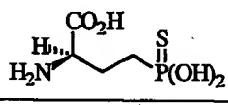
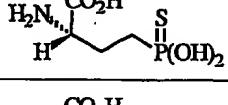
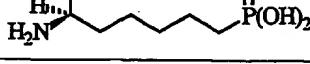
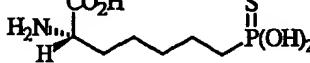
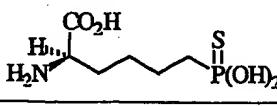
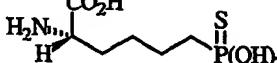
Ethylthiophosphonic acid	$\text{CH}_3\text{CH}_2\overset{\text{S}}{\underset{\parallel}{\text{P}}}(\text{O})_2\text{H}_2$
Methylthiophosphonic acid	$\text{CH}_3\overset{\text{S}}{\underset{\parallel}{\text{P}}}(\text{O})_2\text{H}_2$
<i>tert</i> -Butylthiophosphonic acid	$(\text{CH}_3)_3\overset{\text{S}}{\underset{\parallel}{\text{CPO}}}(\text{O})_2\text{H}_2$
3-Thiophosphonopropanoic acid	$\text{HO}_2\text{CCH}_2\text{CH}_2\overset{\text{S}}{\underset{\parallel}{\text{P}}}(\text{O})_2\text{H}_2$
Phenylthiophosphonic acid	
3-Aminopropylthiophosphonic acid	$\text{NH}_2\text{CH}_2\text{CH}_2\text{CH}_2\overset{\text{S}}{\underset{\parallel}{\text{P}}}(\text{O})_2\text{H}_2$
(<i>dL</i>)-2-Amino-3-thiophosphonopropanoic acid	$\text{HO}_2\text{CCH}(\text{NH}_2)\overset{\text{S}}{\underset{\parallel}{\text{CH}}}(\text{O})_2$
(1-Aminopropyl)thiophosphonic acid	$\text{CH}_3\text{CH}_2\overset{\text{NH}_2}{\underset{\parallel}{\text{CH}}}\overset{\text{S}}{\underset{\parallel}{\text{P}}}(\text{O})_2\text{H}_2$
(<i>dL</i>)-2-Amino-5-thiophosphonopentanoic acid	$\text{HO}_2\text{CCH}(\text{NH}_2)\overset{\text{S}}{\underset{\parallel}{\text{CH}_2\text{CH}_2\text{CH}_2}}\overset{\text{S}}{\underset{\parallel}{\text{P}}}(\text{O})_2\text{H}_2$
(<i>S</i>)-2-Amino-2-methyl-4-thiophosphonobutanoic acid	
D-2-Amino-4-thiophosphonobutanoic acid	
L-2-Amino-4-thiophosphonobutanoic acid	
D-2-Amino-7-thiophosphonoheptanoic acid	
L-2-Amino-7-thiophosphonoheptanoic acid	
D-2-Amino-6-thiophosphonohexanoic acid	
L-2-Amino-6-thiophosphonohexanoic acid	

Table III

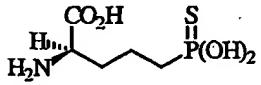
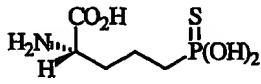
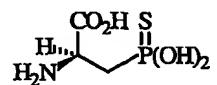
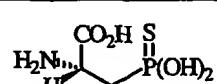
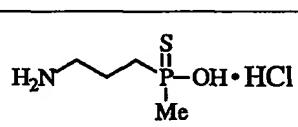
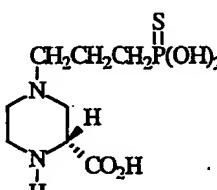
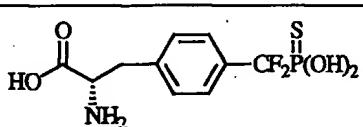
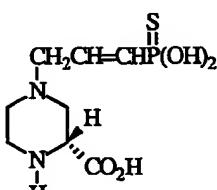
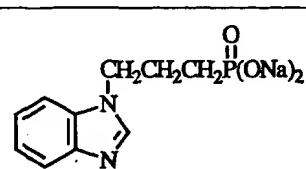
D-2-Amino-5-thiophosphonopentanoic acid	
L-2-Amino-5-thiophosphonopentanoic acid	
D-2-Amino-3-thiophosphonopropanoic acid	
L-2-Amino-3-thiophosphonopropanoic acid	
3-Aminopropyl(methyl)thiophosphinic acid, hydrochloride	
(R)-3-(3-Carboxy-1-piperazinyl)-1-propyl-thiophosphonic acid	
L-4-[Difluoro(thiophosphono)methyl)]-phenylalanine	
(R,E)-4-(3-Thiophosphonoprop-2-enyl)piperazine-2-carboxylic acid	
4-Amino-1-butylphosphonic acid, disodium salt	
4-Amino-1-butylthiophosphonic acid, disodium salt	
1-(3-Phosphonopropyl)-benzimidazole, disodium salt	

Table III

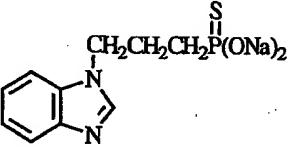
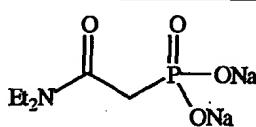
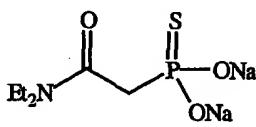
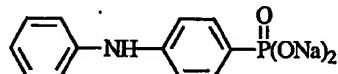
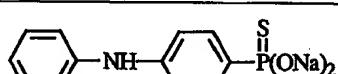
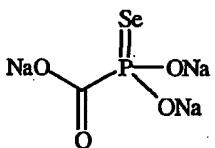
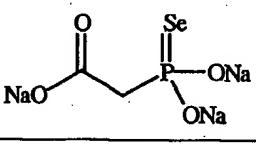
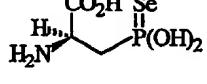
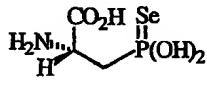
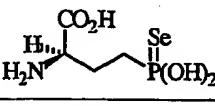
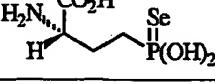
1-(3-Thiophosphonopropyl)-benzimidazole, disodium salt	
3-Dimethylamino-1-propylphosphonic acid, disodium salt	
N,N-Diethylphosphonoacetamide, disodium salt	
N,N-Diethylthiophosphonoacetamide, disodium salt	
Diphenylamine-4-phosphonic acid, disodium salt	
Diphenylamine-4-thiophosphonic acid, disodium salt	
Selenophosphonoformic acid, trisodium salt	
Selenophosphonoacetic acid, trisodium salt	
D-2-Amino-3-selenophosphonopropanoic acid	
L-2-Amino-3-selenophosphonopropanoic acid	
D-2-Amino-4-selenophosphonobutanoic acid	
L-2-Amino-4-selenophosphonobutanoic acid	

Table III

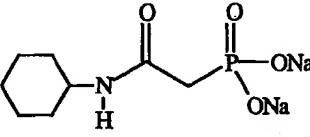
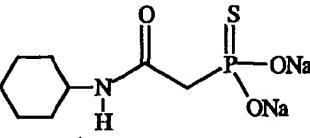
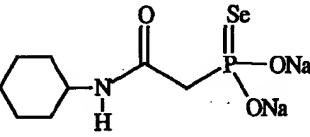
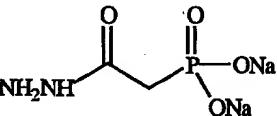
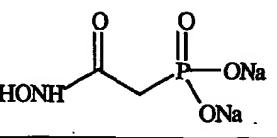
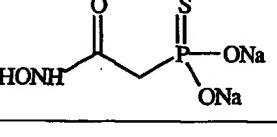
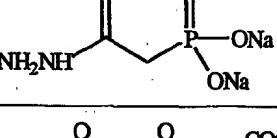
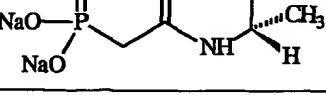
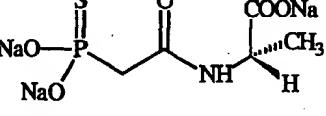
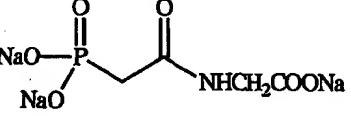
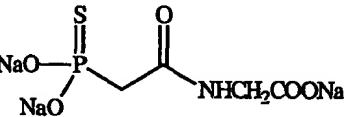
<i>N</i> -Cyclohexylphosphonoacetamide, disodium salt	
<i>N</i> -Cyclohexylthiophosphonoacetamide, disodium salt	
<i>N</i> -Cyclohexylselenophosphonoacetamide, disodium salt	
Phosphonoacetic hydrazide, disodium salt	
<i>N</i> -Hydroxyphosphonoacetamide, disodium salt	
<i>N</i> -Hydroxythiophosphonoacetamide, disodium salt	
Thiophosphonoacetic hydrazide, disodium salt	
<i>N</i> -Phosphonoacetyl-L-alanine, trisodium salt	
<i>N</i> -Thiophosphonoacetyl-L-alanine, trisodium salt	
<i>N</i> -Phosphonoacetyl-Glycine, trisodium salt	
<i>N</i> -Thiophosphonoacetyl-Glycine, trisodium salt	

Table III

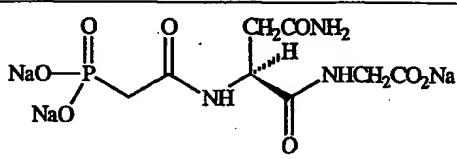
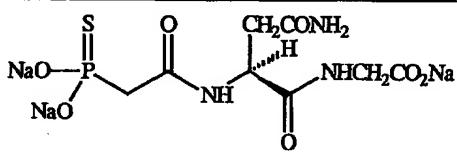
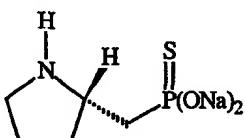
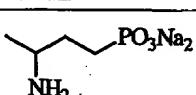
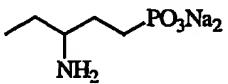
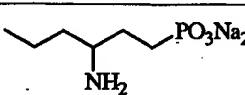
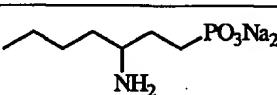
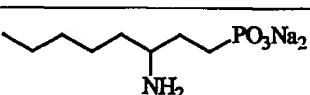
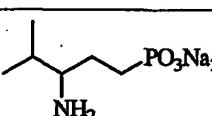
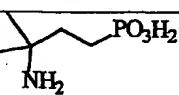
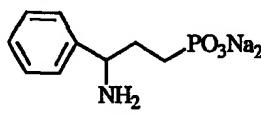
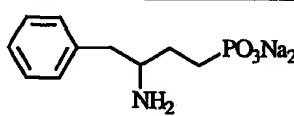
<i>N</i> -(Phosphonoacetyl)-L-asparagine-Glycine, tetrasodium salt	
<i>N</i> -(Thiophosphonoacetyl)-L-asparagine-Glycine, tetrasodium salt	
(S)-2-Pyrrolidinemethylthiophosphonic acid, disodium salt	
(<i>d</i> , <i>l</i>)-3-Amino-butylphosphonic acid, disodium salt	
(<i>d</i> , <i>l</i>)-3-Amino-pentylphosphonic acid, disodium salt	
(<i>d</i> , <i>l</i>)-3-Amino-hexylphosphonic acid, disodium salt	
(<i>d</i> , <i>l</i>)-3-Amino-heptylphosphonic acid, disodium salt	
(<i>d</i> , <i>l</i>)-3-Amino-octylphosphonic acid, disodium salt	
(<i>d</i> , <i>l</i>)-3-Amino-4-methyl-pentylphosphonic acid, disodium salt	
3-Amino-3-methyl-butylphosphonic acid, disodium salt	
(<i>d</i> , <i>l</i>)-3-Amino-3-phenyl-propylphosphonic acid, disodium salt	
(<i>d</i> , <i>l</i>)-3-Amino-4-phenyl-butylphosphonic acid, disodium salt	

Table III

(<i>D,L</i>)-3-Amino-4-phenyl-pentyphosphonic acid, disodium salt	
(<i>D,L</i>)-3-Amino-3-phenyl-butylphosphonic acid, disodium salt	
(<i>D,L</i>)-2-Amino-2-(2-phosphonoethyl)-1,2,3,4-tetrahydronaphthalene, disodium salt	
1-Amino-1-(2-phosphonoethyl)-cyclohexane, disodium salt	
(<i>D,L</i>)-2-(2-Amino-4-phosphonobutoxy)tetrahydropyran, disodium salt	
(<i>D,L</i>)-3-Amino-4-hydroxybutylphosphonic acid, disodium salt	
3-Phosphonopropanesulfonic acid, trisodium salt	
Pamidronic acid (3-Amino-1-hydroxypropane-1,1-bisphosphonic acid)	
3-Amino-1-hydroxypropane-1,1-bisphosphonic acid, tetrasodium salt	
Diethyl 2-pyrrolidinylphosphonate	
2-Pyrrolidinylphosphonic acid, disodium salt	
1,1-Dioxo-2-(3-phosphonopropyl)-isothiazoline, disodium salt	
1,1-Dioxo-2-(3-thiophosphonopropyl)-isothiazolidin, disodium salt	

Table III

2-Deoxy-2-phosphonoacetylamino-D-glucose	
2-Deoxy-2-thiophosphonoacetylamino-D-glucose	
1-Amino-3-sulfopropane-1,1-bisphosphonic acid	
1-Amino-3-sulfopropane-1,1-bisphosphonic acid, pentasodium salt	
3-Hydroxy-3-(2-pyridyl)propenyl-2-phosphonic acid, disodium salt	
3-Hydroxy-3-(3-pyridyl)propenyl-2-phosphonic acid, disodium salt	
3-Hydroxy-3-(4-pyridyl)propenyl-2-phosphonic acid, disodium salt	
3-Amino-3-(2-pyridyl)propenyl-2-phosphonic acid, disodium salt	
3-Amino-3-(3-pyridyl)propenyl-2-phosphonic acid, disodium salt	

Table III

3-Amino-3-(4-pyridyl)propenyl-2-phosphonic acid, disodium salt	
1,4-Diamino-1-(3-pyridyl)butyl-2-phosphonic acid, disodium salt	
1,4-Diamino-4-methyl-1-(3-pyridyl)pentyl-2-phosphonic acid, disodium salt	
1,4-Diamino-4-methyl-1-(2-pyridyl)pentyl-2-phosphonic acid, disodium salt	
1,4-Diamino-4-methyl-1-(4-pyridyl)pentyl-2-phosphonic acid, disodium salt	
1,3-Diaminopropane-1,1-bisphosphonic acid, tetrasodium salt	
1-Amino-3-dimethylaminopropane-1,1-bisphosphonic acid, tetrasodium salt	
3-Dimethylamino-1-hydroxypropane-1,1-bisphosphonic acid, tetrasodium salt	
1-Hydroxy-3-(methylphenylamino)-propane-1,1-bisphosphonic acid, tetrasodium salt	
1-Amino-3-(methylphenylamino)propane-1,1-bisphosphonic acid, tetrasodium salt	

Table III

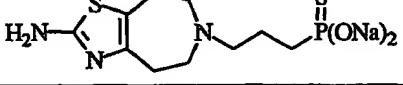
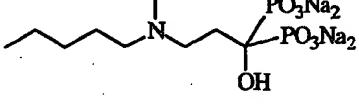
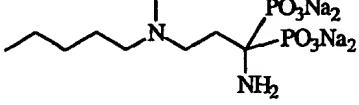
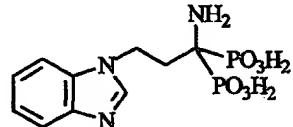
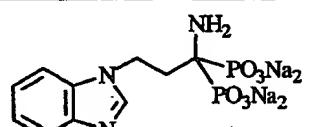
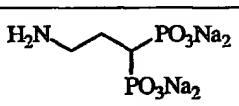
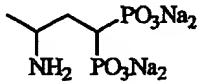
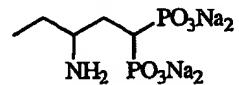
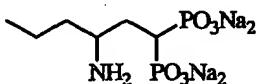
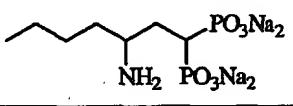
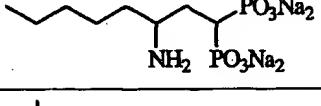
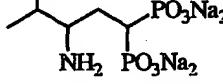
3-(2-Amino-4,5,7,8-tetrahydro-6 <i>H</i> -thiazolo[4,5- <i>d</i>]azepin-6-yl)propyl-phosphonic acid, disodium salt	
Ibandronic acid, tetrasodium salt (1-Hydroxy-3-(methylpentylamino)-propane-1,1-bisphosphonic acid, tetrasodium salt)	
1-Amino-3-(methylpentylamino)propane-1,1-bisphosphonic acid, tetrasodium salt	
1-Amino-3-(1-benzimidazolyl)propane-1,1-bisphosphonic acid	
1-Amino-3-(1-benzimidazolyl)propane-1,1-bisphosphonic acid, tetrasodium salt	
3-Aminopropane-1,1-bisphosphonic acid, tetrasodium salt	
(<i>D,L</i>)-3-Aminobutane-1,1-bisphosphonic acid, tetrasodium salt	
(<i>D,L</i>)-3-Aminopentane-1,1-bisphosphonic acid, tetrasodium salt	
(<i>D,L</i>)-3-Aminohexane-1,1-bisphosphonic acid, tetrasodium salt	
(<i>D,L</i>)-3-Aminoheptane-1,1-bisphosphonic acid, tetrasodium salt	
(<i>D,L</i>)-3-Aminooctane-1,1-bisphosphonic acid, tetrasodium salt	
(<i>D,L</i>)-3-Amino-4-methylpentane-1,1-bisphosphonic acid, tetrasodium salt	

Table III

3-Amino-3-methylbutane-1,1-bisphosphonic acid, tetrasodium salt	
(<i>dL</i>)-3-Amino-3-phenylpropane-1,1-bisphosphonic acid, tetrasodium salt	
(<i>dL</i>)-3-Amino-4-phenylbutane-1,1-bisphosphonic acid, tetrasodium salt	
3-Amino-4-phenylpentane-1,1-bisphosphonic acid, tetrasodium salt	
(<i>dL</i>)-3-Amino-3-phenylbutane-1,1-bisphosphonic acid, tetrasodium salt	
(<i>dL</i>)-2-(2-Amino-1,2,3,4-tetrahydronaphthalenyl)ethane-1,1-bisphosphonic acid, tetrasodium salt	
2-(1-Aminocyclohexyl)ethane-1,1-bisphosphonic acid, tetrasodium salt	
2-(2-Amino-4,4-bisphosphonobutoxy)-tetrahydropyran, tetrasodium salt	
(<i>dL</i>)-3-Amino-4-hydroxybutane-1,1-bisphosphonic acid, tetrasodium salt	
(<i>S</i>)-Hydroxy(2-pyrrolidinyl)methane-bisphosphonic acid, sodium salt	
Hydroxy[(2 <i>S</i> ,4 <i>R</i>)-4-hydroxy-2-pyrrolidinyl]methanebisphosphonic acid tetrasodium salt	

Table III

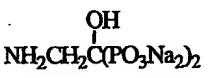
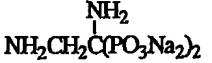
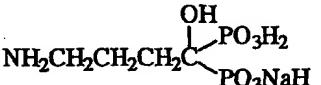
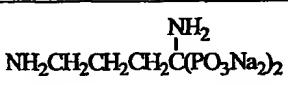
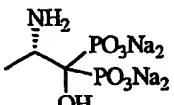
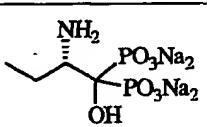
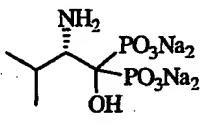
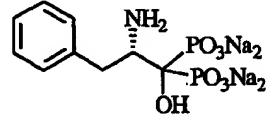
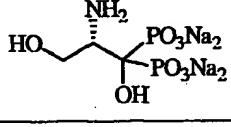
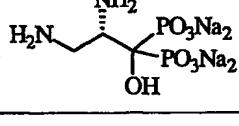
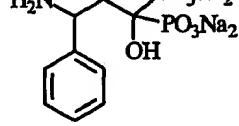
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1,4-Diaminobutane-1,1-bisphosphonic acid, tetrasodium salt	
5-Amino-1-hydroxypentane-1,1-bisphosphonic acid, tetrasodium salt	
1,5-Diaminopentane-1,1-bisphosphonic acid, tetrasodium salt	
(S)-2-Amino-1-hydroxypropane-1,1-bisphosphonic acid, tetrasodium salt	
(S)-2-Amino-1-hydroxybutane-1,1-bisphosphonic acid, tetrasodium salt	
(S)-2-Amino-1-hydroxy-3-methylbutane-1,1-bisphosphonic acid, tetrasodium salt	
(S)-2-Amino-1-hydroxy-3-phenylpropane-1,1-bisphosphonic acid, tetrasodium salt	
(S)-2-Amino-1,3-dihydroxypropane-1,1-bisphosphonic acid, tetrasodium salt	
(S)-2,3-Diamino-1-hydroxypropane-1,1-bisphosphonic acid, tetrasodium salt	
(dl)-3-Amino-1-hydroxy-3-phenylpropane-1,1-bisphosphonic acid, tetrasodium salt	

Table III

(S)-3-Amino-2-(4-chlorophenyl)-1-hydroxypropane-1,1-bisphosphonic acid, tetrasodium salt	
(S)-2-Amino-3-(4-aminophenyl)-1-hydroxypropane-1,1-bisphosphonic acid, tetrasodium salt	
N-Phosphonomethylglycine	
N-Phosphonomethylglycine, trisodium salt	
2-Phosphonomethylglutaric acid, tetrasodium salt	
2-Phosphonomethylsuccinic acid, tetrasodium salt	
(2R,4S)-4-Phosphonomethylpipecolinic acid, trisodium salt	
(2R,4S)-4-Phosphonomethylpipecolinamide, disodium salt	
N-Phosphonomethylglycine	
N-Phosphonomethylglycine, trisodium salt	
3-[6-Methoxy-2-(1,2,3,4-tetrahydroisoquinolinyl)]propylphosphonic acid, disodium salt	

Table III

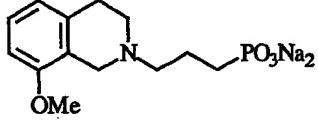
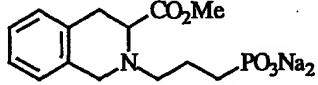
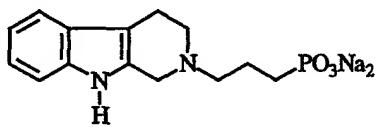
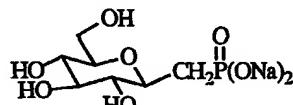
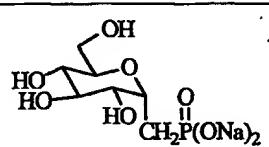
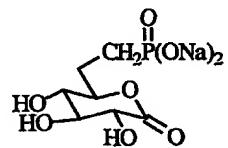
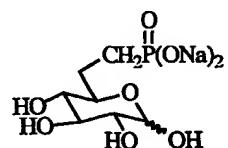
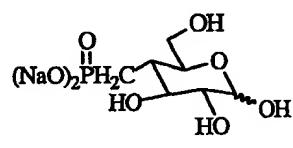
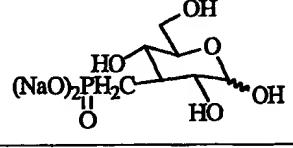
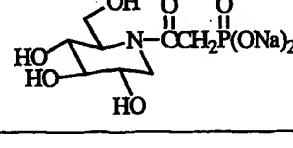
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3-[2-(3-Methoxycarbonyl-1,2,3,4-tetrahydroisoquinolinyl)]-propylphosphonic acid disodium salt	
2-(3-Phosphonopropyl)-1,2,3,4-tetrahydro-9 <i>H</i> -pyrido[3,4- <i>b</i>]indole, disodium salt	
β -D-Glucopyranosylmethylphosphonic acid, disodium salt	
α -D-Glucopyranosylmethylphosphonic acid, disodium salt	
6-Deoxy-6-C-phosphonomethyl-D-glucono- δ -lactone, disodium salt	
6-Deoxy-6-C-phosphonomethyl-D-glucose, disodium salt	
4-Deoxy-4-C-phosphonomethyl-D-glucose, disodium salt	
3-Deoxy-3-C-phosphonomethyl-D-glucose, disodium salt	
1-Deoxy-N-phosphonoacetylinojirimycin, disodium salt	

Table III

(1,5-Dideoxy-1,5-imino- α -D-glucopyranosyl)methylphosphonic acid, disodium salt	
1,6-Dideoxy-6-C-phosphonomethyl-nojirimycin, disodium salt	

Table IV

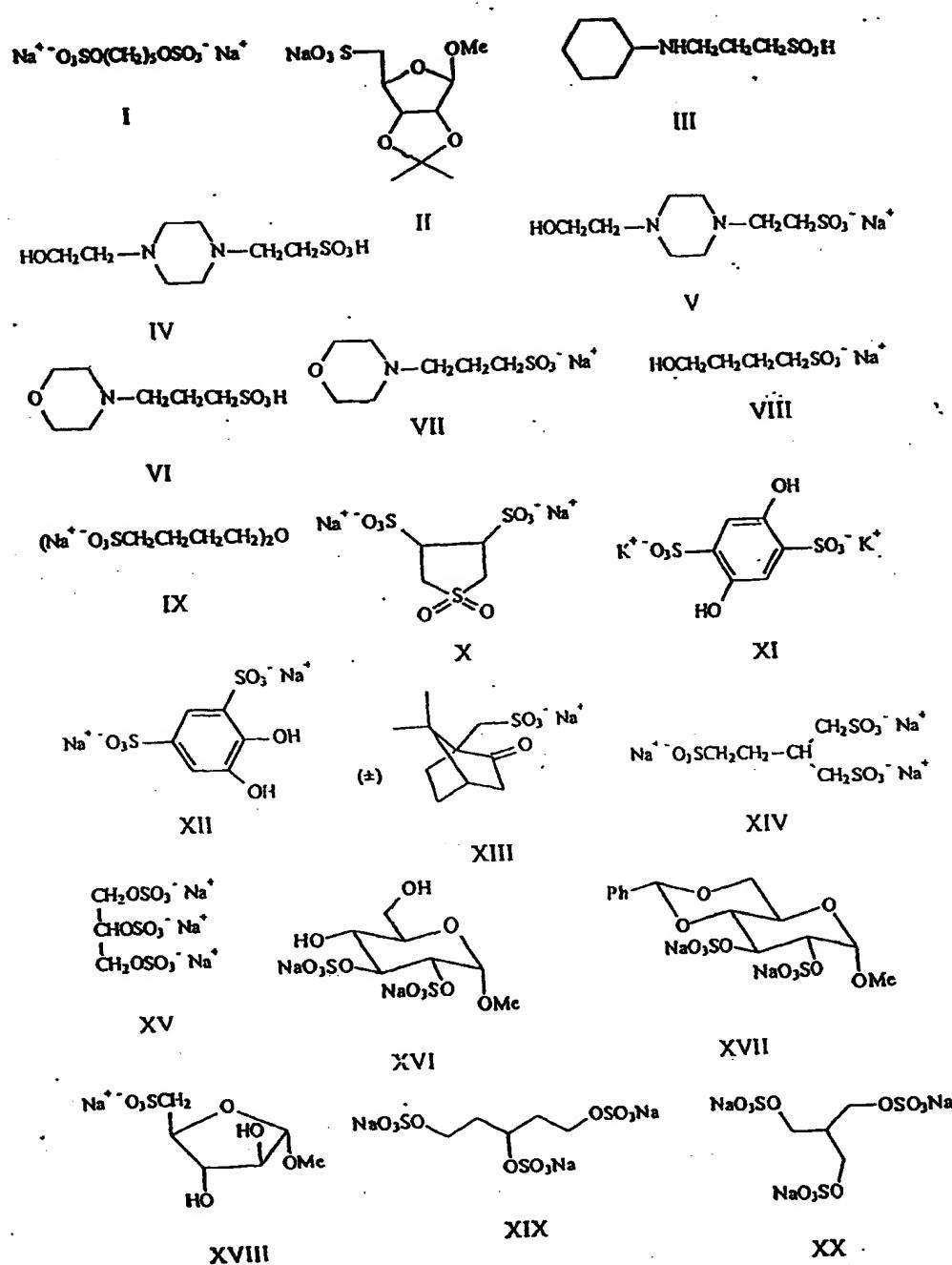


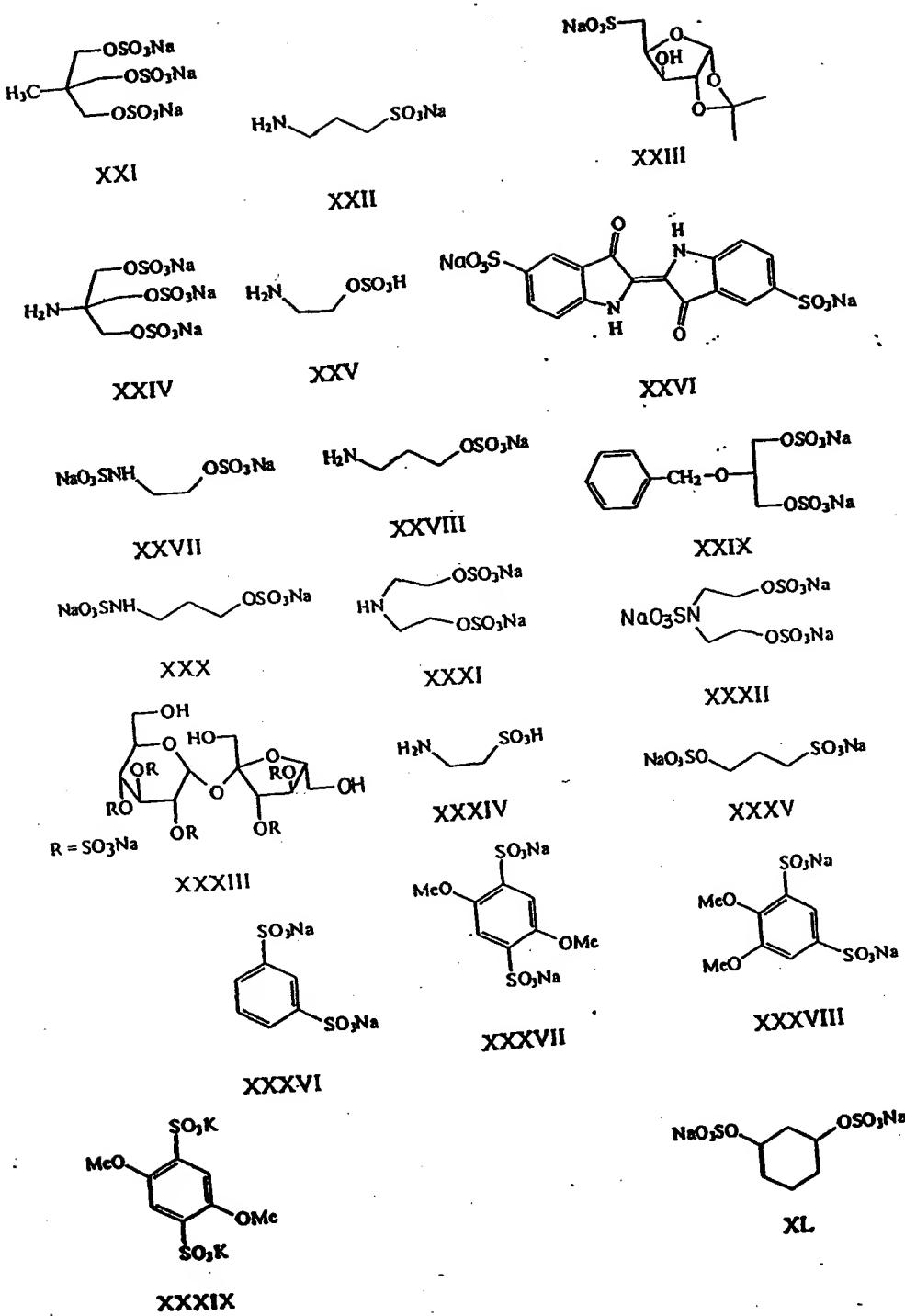
Table IV

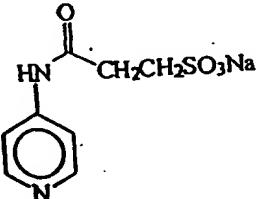
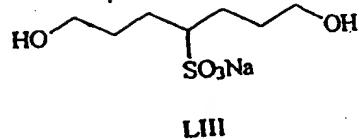
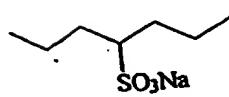
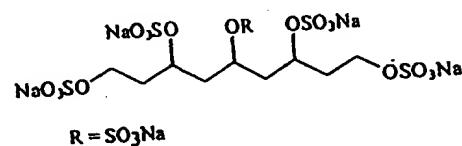
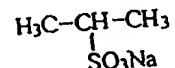
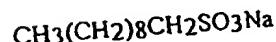
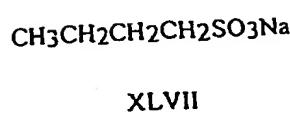
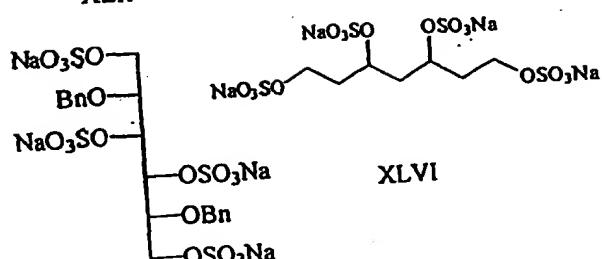
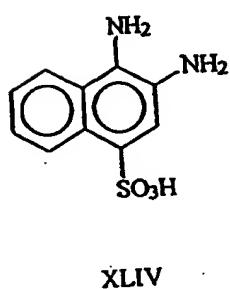
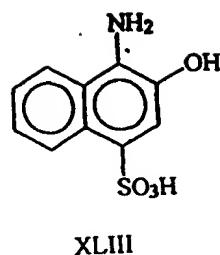
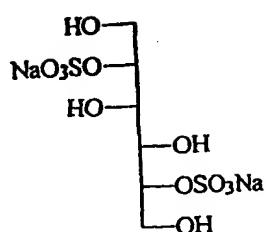
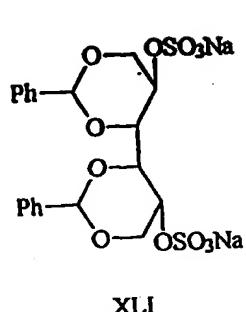
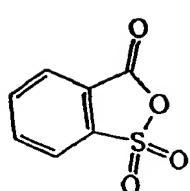
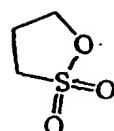
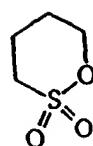
Table IV

Table IV

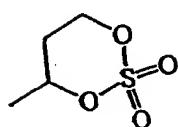
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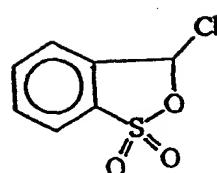
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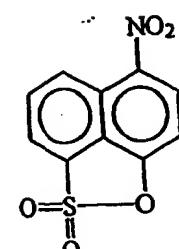
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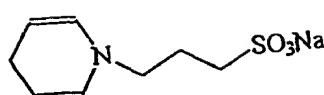
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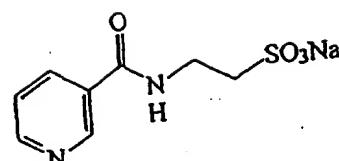
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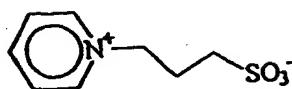
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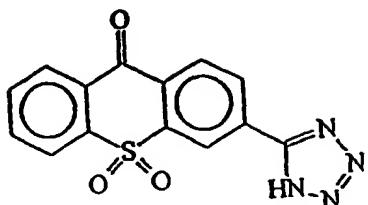
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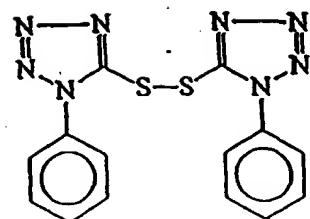
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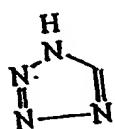
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Table IV

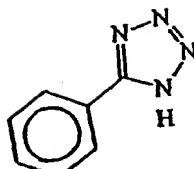
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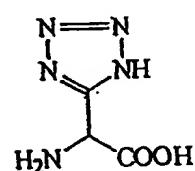
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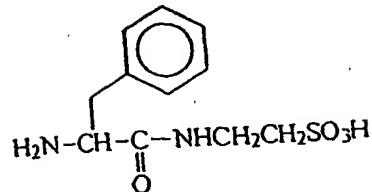
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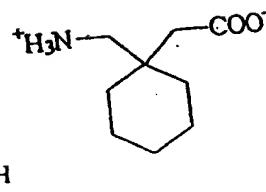
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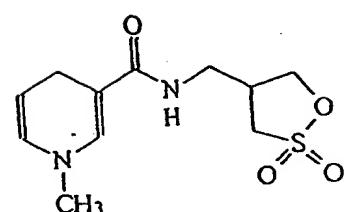
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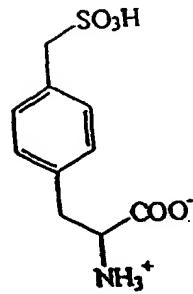
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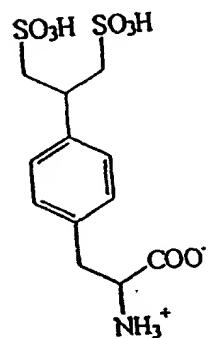
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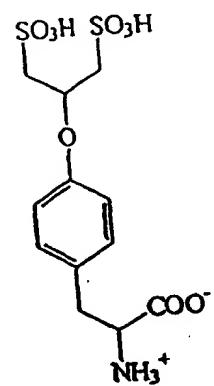
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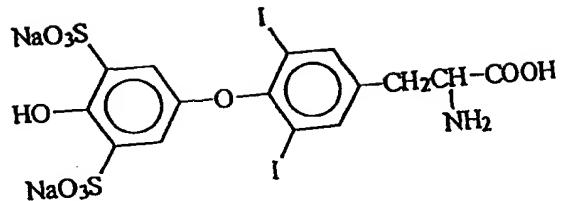
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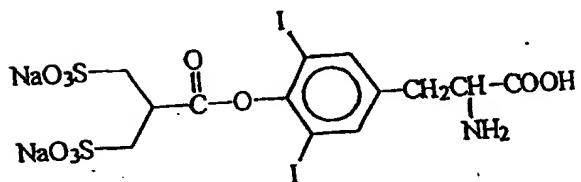
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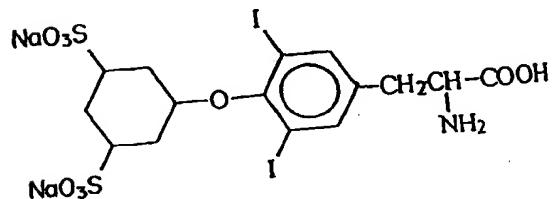
LXXIV

Table IV

LXXV



LXXVI



LXXVII

EQUIVALENTS

Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, numerous equivalents to the specific procedures described herein. Such equivalents are considered to be within the scope of the present invention and are covered by 5 the following claims. The contents of all references, issued patents, and published patent applications cited throughout this application are hereby incorporated by reference. The appropriate components, processes, and methods of those patents, applications and other documents may be selected for the present invention and embodiments thereof.

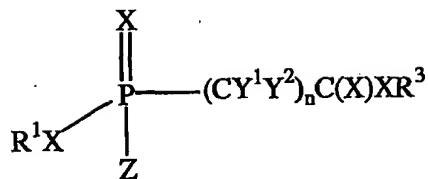
CLAIMS

What is claimed is:

1. A method of inhibiting cerebral amyloid angiopathy, comprising contacting a blood vessel wall cell with an A β 40 inhibitor, such that cerebral amyloid angiopathy is inhibited.
- 5 2. The method of claim 1 wherein the A β 40 inhibitor has the following structure:



3. The method of claim 1, wherein said A β 40 inhibitor is selected from the group consisting of ethanesulfonic acid, 1,2-ethanedisulfonic acid, 1-propanesulfonic acid, 1,3-propanedisulfonic acid, 1,4-butanedisulfonic acid, 1,5-pentanedisulfonic acid, 2-aminoethanesulfonic acid, 4-hydroxy-1-butanesulfonic acid, and pharmaceutically acceptable salts thereof.
- 10 4. The method of claim 1, wherein said A β 40 inhibitor is selected from the group consisting of 1-butanesulfonic acid, 1-decanesulfonic acid, 2-propanesulfonic acid, 3-pentanesulfonic acid, 4-heptanesulfonic acid, and pharmaceutically acceptable salts thereof.
- 15 5. The method of claim 1, wherein said A β 40 inhibitor is 1,7-dihydroxy-4-heptanesulfonic acid, or a pharmaceutically acceptable salt thereof.
6. The method of claim 1, wherein said A β 40 inhibitor is 3-amino-1-propanesulfonic acid, or a salt thereof.
7. The method of claim 1, wherein said blood vessel wall cell is selected from the group consisting of blood vessel wall smooth muscle cells, pericytes and endothelial cells.
- 20 8. The method of claim 1, wherein said blood vessel wall cell is a blood vessel wall smooth muscle cell.
9. The method of claim 1 wherein the death of said blood vessel wall cell is prevented.
10. The method of claim 1 wherein structural changes to said blood vessel wall cell are prevented.
- 25 11. The method of claim 1, wherein said A β 40 inhibitor is a peptide or a peptidomimetic which interacts with specific regions of the A β peptide.
12. The method of claim 1, wherein said A β 40 inhibitor has the following structure:



wherein

Z is XR² or R⁴;

R¹ and R² are each independently hydrogen, a substituted or unsubstituted aliphatic group, an aryl group, a heterocyclic group, or a salt-forming cation;

R³ is hydrogen, lower alkyl, aryl, or a salt-forming cation;

R⁴ is hydrogen, lower alkyl, aryl or amino;

X is, independently for each occurrence, O or S;

Y¹ and Y² are each independently hydrogen, halogen, alkyl, amino, hydroxy, alkoxy, or aryloxy; and

n is an integer from 0 to 12.

13. The method of claim 1, wherein said Aβ40 inhibitor is administered in a pharmaceutically acceptable formulation.

14. The method of claim 13, wherein said pharmaceutically acceptable formulation is a dispersion system.

15. The method of claim 14, wherein said pharmaceutically acceptable formulation comprises a lipid-based formulation.

16. The method of claim 15, wherein said pharmaceutically acceptable formulation comprises a liposome formulation.

20 17. The method of claim 16, wherein said pharmaceutically acceptable formulation comprises a multivesicular liposome formulation.

18. The method of claim 15, wherein said pharmaceutically acceptable formulation comprises a polymeric matrix.

19. The method of claim 18, wherein said polymeric matrix is selected from the group consisting of naturally derived polymers, such as albumin, alginate, cellulose derivatives, collagen, fibrin, gelatin, and polysaccharides.

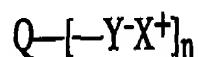
20. The method of claim 18, wherein said polymeric matrix is selected from the group consisting of synthetic polymers such as polyesters (PLA, PLGA), polyethylene glycol, poloxomers, polyanhydrides, and pluronic.

21. The method of claim 18, wherein said polymeric matrix is in the form of microspheres.

5 22. The method of claim 13, wherein the pharmaceutically acceptable formulation provides sustained delivery of said A_β40 inhibitor to a subject.

23. A method of treating a disease state characterized by cerebral amyloid angiopathy in a subject, comprising administering an A_β40 inhibitor to said subject, such that said disease state characterized by cerebral amyloid angiopathy is treated.

10 24. The method of claim 23 wherein said A_β40 inhibitor has the structure:



wherein Y⁻ is an anionic group at physiological pH; Q is a carrier group; X⁺ is a cationic group; and n is an integer selected such that the biodistribution of the A_β40 inhibitor for an intended target site is not prevented while maintaining activity of the A_β40 inhibitor, such that cerebral amyloid angiopathy is inhibited.

15 25. The method of claim 23, wherein said A_β40 inhibitor is selected from the group consisting of ethanesulfonic acid, 1,2-ethanedisulfonic acid, 1-propanesulfonic acid, 1,3-propanedisulfonic acid, 1,4-butanedisulfonic acid, 1,5-pentanedisulfonic acid, 2-aminoethanesulfonic acid, 4-hydroxy-1-butanesulfonic acid, and pharmaceutically acceptable salts thereof.

29. The method of claim 23, wherein said blood vessel wall cell is selected from the group consisting of blood vessel wall smooth muscle cells, pericytes and endothelial cells.

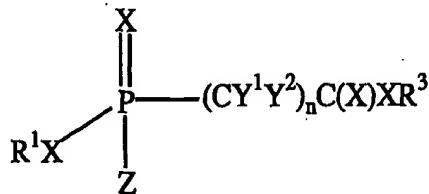
30. The method of claim 23, wherein said blood vessel wall cell is a blood vessel wall smooth muscle cell.

5 31. The method of claim 23 wherein the death of said blood vessel wall cell is prevented.

32. The method of claim 23 wherein structural changes to said blood vessel wall cell are prevented.

33. The method of claim 23, wherein said A_β40 inhibitor is a peptide or a peptidomimetic which interacts with specific regions of the A_β peptide.

10 34. The method of claim 23, wherein said A_β40 inhibitor has the following structure:



wherein

Z is XR² or R⁴;

R¹ and R² are each independently hydrogen, a substituted or unsubstituted

15 aliphatic group, an aryl group, a heterocyclic group, or a salt-forming cation;

R³ is hydrogen, lower alkyl, aryl, or a salt-forming cation;

R⁴ is hydrogen, lower alkyl, aryl or amino;

X is, independently for each occurrence, O or S;

Y¹ and Y² are each independently hydrogen, halogen, alkyl, amino, hydroxy,

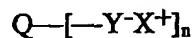
alkoxy, or aryloxy; and

20 n is an integer from 0 to 12.

35. The method of claim 23, wherein said A_β40 inhibitor is administered in a pharmaceutically acceptable formulation.

36. The method of claim 35, wherein said pharmaceutically acceptable formulation is a dispersion system.

37. The method of claim 35, wherein said pharmaceutically acceptable formulation comprises a lipid-based formulation.
38. The method of claim 35, wherein said pharmaceutically acceptable formulation comprises a liposome formulation.
- 5 39. The method of claim 38, wherein said pharmaceutically acceptable formulation comprises a multivesicular liposome formulation.
40. The method of claim 35, wherein said pharmaceutically acceptable formulation comprises a polymeric matrix.
- 10 41. The method of claim 41, wherein said polymeric matrix is selected from the group consisting of naturally derived polymers, such as albumin, alginate, cellulose derivatives, collagen, fibrin, gelatin, and polysaccharides.
42. The method of claim 41, wherein said polymeric matrix is selected from the group consisting of synthetic polymers such as polyesters (PLA, PLGA), polyethylene glycol, poloxomers, polyanhydrides, and pluronic.
- 15 43. The method of claim 41, wherein said polymeric matrix is in the form of microspheres.
44. The method of claim 35, wherein the pharmaceutically acceptable formulation provides sustained delivery of said A_β40 inhibitor to a subject.
45. A method of inhibiting cerebral amyloid angiopathy in a subject, comprising administering an A_β40 inhibitor to said patient in an effective amount and manner such that said A_β40
20 inhibitor contacts a blood vessel wall cell in said patient and that cerebral amyloid angiopathy is inhibited.
46. The method of claim 45 wherein the A_β40 inhibitor has the following structure:



wherein Y⁻ is an anionic group at physiological pH; Q is a carrier group; X⁺ is a cationic group; and n is an integer selected such that the biodistribution of the A_β40 inhibitor for an intended target site is not prevented while maintaining activity of the A_β40 inhibitor, such that cerebral amyloid angiopathy is inhibited.

47. The method of claim 45, wherein said A β 40 inhibitor is selected from the group consisting of ethanesulfonic acid, 1,2-ethanedisulfonic acid, 1-propanesulfonic acid, 1,3-propanedisulfonic acid, 1,4-butanedisulfonic acid, 1,5-pentanedisulfonic acid, 2-aminoethanesulfonic acid, 4-hydroxy-1-butanesulfonic acid, and pharmaceutically acceptable salts thereof.

5 48. The method of claim 45, wherein said A β 40 inhibitor is selected from the group consisting of 1-butanesulfonic acid, 1-decanesulfonic acid, 2-propanesulfonic acid, 3-pentanesulfonic acid, 4-heptanesulfonic acid, and pharmaceutically acceptable salts thereof.

10 49. The method of claim 45, wherein said A β 40 inhibitor is 1,7-dihydroxy-4-heptanesulfonic acid, or a pharmaceutically acceptable salt thereof.

50. The method of claim 45, wherein said A β 40 inhibitor is 3-amino-1-propanesulfonic acid, or a salt thereof.

51. The method of claim 45, wherein said blood vessel wall cell is selected from the group consisting of blood vessel wall smooth muscle cells, pericytes and endothelial cells.

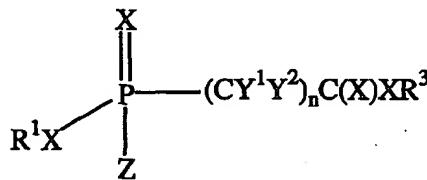
15 52. The method of claim 45, wherein said blood vessel wall cell is a blood vessel wall smooth muscle cell.

53. The method of claim 45 wherein the death of said blood vessel wall cell is prevented.

54. The method of claim 45 wherein structural changes to said blood vessel wall cell are prevented.

20 55. The method of claim 45, wherein said A β 40 inhibitor is a peptide or a peptidomimetic which interacts with specific regions of the A β peptide.

56. The method of claim 45, wherein said A β 40 inhibitor has the following structure:



wherein

25 Z is XR² or R⁴;

R¹ and R² are each independently hydrogen, a substituted or unsubstituted aliphatic group, an aryl group, a heterocyclic group, or a salt-forming cation; R³ is hydrogen, lower alkyl, aryl, or a salt-forming cation; R⁴ is hydrogen, lower alkyl, aryl or amino;

5 X is, independently for each occurrence, O or S;

Y¹ and Y² are each independently hydrogen, halogen, alkyl, amino, hydroxy, alkoxy, or aryloxy; and

n is an integer from 0 to 12.

57. The method of claim 45, wherein said Aβ40 inhibitor is administered in a pharmaceutically acceptable formulation.

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58. The method of claim 57, wherein said pharmaceutically acceptable formulation is a dispersion system.

59. The method of claim 58, wherein said pharmaceutically acceptable formulation comprises a lipid-based formulation.

15 60. The method of claim 59, wherein said pharmaceutically acceptable formulation comprises a liposome formulation.

61. The method of claim 59, wherein said pharmaceutically acceptable formulation comprises a multivesicular liposome formulation.

62. The method of claim 57, wherein said pharmaceutically acceptable formulation comprises a 20 polymeric matrix.

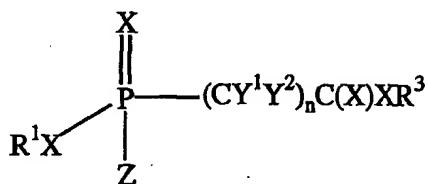
63. The method of claim 62, wherein said polymeric matrix is selected from the group consisting of naturally derived polymers, such as albumin, alginate, cellulose derivatives, collagen, fibrin, gelatin, and polysaccharides.

64. The method of claim 62, wherein said polymeric matrix is selected from the group 25 consisting of synthetic polymers such as polyesters (PLA, PLGA), polyethylene glycol, poloxomers, polyanhydrides, and pluronic.

65. The method of claim 62, wherein said polymeric matrix is in the form of microspheres.

66. The method of claim 57, wherein the pharmaceutically acceptable formulation provides sustained delivery of said Aβ40 inhibitor to a subject.

67. A method of inhibiting cerebral amyloid angiopathy, comprising contacting a blood vessel wall cell with a A β 40 inhibitor having the structure:



wherein

5 Z is XR² or R⁴;

R¹ and R² are each independently hydrogen, a substituted or unsubstituted aliphatic group, an aryl group, a heterocyclic group, or a salt-forming cation;

R³ is hydrogen, lower alkyl, aryl, or a salt-forming cation;

R⁴ is hydrogen, lower alkyl, aryl or amino;

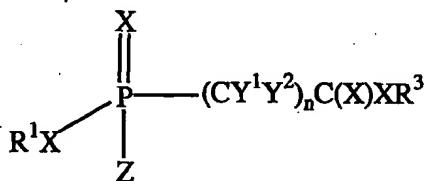
10 X is, independently for each occurrence, O or S;

Y¹ and Y² are each independently hydrogen, halogen, alkyl, amino, hydroxy, alkoxy, or aryloxy; and

n is an integer from 0 to 12, such that cerebral amyloid angiopathy is inhibited.

68. A method of inhibiting cerebral amyloid angiopathy in a subject, comprising administering

15 an A β 40 inhibitor to said patient in an effective amount and manner such that said A β 40 inhibitor contacts a blood vessel wall cell in said patient, said A β 40 inhibitor having the structure:



wherein

20 Z is XR² or R⁴;

R¹ and R² are each independently hydrogen, a substituted or unsubstituted aliphatic group, an aryl group, a heterocyclic group, or a salt-forming cation;

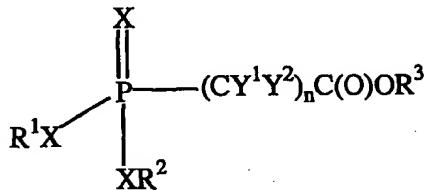
R³ is hydrogen, lower alkyl, aryl, or a salt-forming cation;

R⁴ is hydrogen, lower alkyl, aryl or amino;

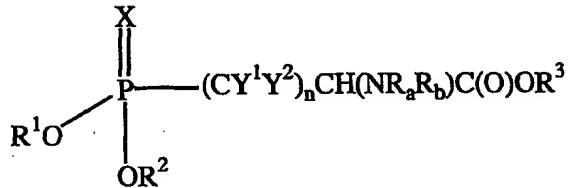
25 X is, independently for each occurrence, O or S;

Y^1 and Y^2 are each independently hydrogen, halogen, alkyl, amino, hydroxy, alkoxy, or aryloxy; and
 n is an integer from 0 to 12,
such that cerebral amyloid angiopathy is inhibited.

5 69. The method of claim 68 wherein said A β 40 inhibitor has the structure:



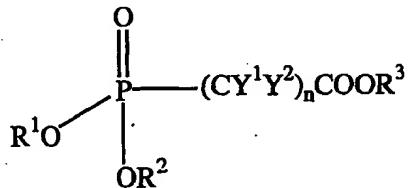
70. The method of claim 68 wherein said A β 40 inhibitor has the structure:



10 wherein R_a and R_b are each independently hydrogen, alkyl, aryl, or heterocyclyl, or R_a and R_b , taken together with the nitrogen atom to which they are attached, form a cyclic moiety having from 3 to 8 atoms in the ring, and n is an integer from 0 to 6.

71. The method of claim 70 wherein R_a and R_b are each hydrogen.

72. The method of claim 68 wherein said A β 40 inhibitor has the structure:

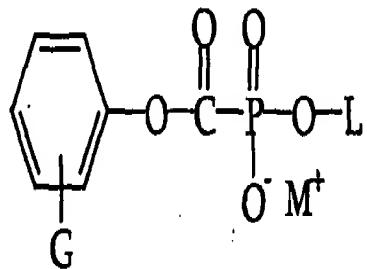


15 wherein R^1 and R^2 are each independently hydrogen, an aliphatic group, an aryl group, a heterocyclic group, or a salt-forming cation; R^3 is hydrogen, lower alkyl, aryl, or a salt-forming cation; Y^1 and Y^2 are each independently hydrogen, halogen (e.g., F, Cl, Br, or I), lower alkyl, hydroxy, alkoxy, or aryloxy; and n is an integer from 0 to 12.

20 73. The method of claim 68 wherein R^1 and R^2 are an aliphatic group selected from the group consisting of a branched or straight-chain aliphatic moiety having from about 1 to 24 carbon

atoms or a branched or straight-chain aliphatic moiety having from about 10-24 carbon atoms, in the chain; and an unsubstituted or substituted cyclic aliphatic moiety having from 4 to 7 carbon atoms in the aliphatic ring.

74. A method of inhibiting cerebral amyloid angiopathy in a subject, comprising administering
5 an A β 40 inhibitor to said patient in an effective amount and manner such that said A β 40 inhibitor contacts a blood vessel wall cell in said patient, said A β 40 inhibitor having the structure:



wherein G represents hydrogen or one or more substituents on the aryl ring and L is a
10 substituted alkyl group, and M $^+$ is a counter ion, such that cerebral amyloid angiopathy is inhibited.

75. The method of claim 74 where G is hydrogen or an electron-donating group.

76. The method of claim 74 where G is an electron-withdrawing group at the meta position.

77. A method of diagnosing cerebral amyloid angiopathy in a subject, comprising administering